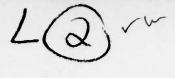
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VOLUME I OF II

RESEARCH IN DRUG DEVELOPMENT AGAINST VIRAL DISEASES OF MILITARY IMPORTANCE (BIOLOGICAL TESTING)

Final Report

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During 1989 at the request of Ms. C. Susan Tiffany (Contract Specialist, Ft. Detrick, Frederick, Maryland) the Adenovirus (AD2) and the Vesicular Stomatitis Virus (VSV) were deleted from the primary screen. Also, all in vivo work was discontinued due to military budget cuts. During 1989 - 1990, antiviral prescreen testing (in vitro) of planextracts against Punta Toro Virus (PT), Yellow Fever Virus (YF) and Venezuelan Equine Encephalitis Virus (VE) was added to the screening program at the rate-of-testing- of approximately 3,000 extracts per year per virus.

During this contract period, a number of compounds were found to exhibit highly significant activity against VV in vitro. Compounds AVS-5569, 5568, 1214, 5219, 0303, 7449 and 6193 demonstrated the greatest in vitro promise, having SI's that ranged from 19 - > 320. The most effective compounds against the Adenovirus were AVS-2296, 2700, 2980, 2986, 3593, 4070 and 4167. Sixty-two excellent lead compound were found against the Yellow Fever Virus in vitro and 45 compounds appeared to have excellent in vitro antiviral potential against the Japanese Encephalitis Virus. Fifteen compounds demonstrated excellent in vitro antiviral activity against Venezuelan Equine Encephalomyelitis Virus. At least 16 excellent lead compounds were found against the Punta Toro Virus in vitro and 24 compounds produced excellent in vitro antiviral activity against Sandfly Fever Virus. AVS-6724 demonstrated broad spectrum activity against YF, JE, SF, PT, VV and VE Viruses. Five compounds were found highly active (therapeutic indices of > 100 against the Pichinde Virus by the plaque-reduction assay in vitro.

The prescreen protocol (YF, JE and VE viruses) has successfully identified potential active materials ($\sim 5\%$) for further confirmatory testing. Confirmatory testing of these potential active compounds were carried out in the primary screen against a broader range of more virulent viruses (VV, YF, JE, VE, PT and SF). Sixty-seven percent of the prescreen compounds showed some degree of activity against one or more of these virulent viruses.

Significant anti-HIV in vitro activity was observed with 14 AVS compounds with therapeutic indices that ranged from > 1000 to > 54. One NCI compound, NSC 614846 demonstrated significant activity comparable to the positive control drug, ddC. Several AVS compounds showed confirmed anti-HIV activity versus the Feline leukemia virus, the Simian and Murine AIDS viruses.

For in vivo antiviral compound evaluations, 2 out of 19 compound had significant activity against the Pichinde Virus in hamsters. Seven compounds had demonstrable activity against JE in mice. Of these seven compounds, AVS-5587 may be the most desirable as the window between the toxicity and efficacy is broader than that seen with the other compounds. Further studies of this compound and its analogs are strongly recommended. AVS-1752 (Ara-A) had demonstrable activity against the Vaccinia Virus in the intracranial challenge model in mice. Nine compounds had some degree of activity against Vaccinia Virus-induced tailpox lesions. The compound with the greatest activity other than the positive control drug (Ara-A) was AVS-3679.

FOREWORD

Citations of commercial organizations and trade names in this report do not constitute an official Department of the Army endorsement or approval of the products or services of these organizations.

In conducting the research described in this report, the investigators adhered to all safety, security, and biocontainment requirements specified in the contract, including all U.S. Public Health Service biosafety guidelines contained in "Biosafety in Microbiological and Biomedical Laboratories", DHHS Publication No. (CDC) 86-23 (March 1985).



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TABLE OF CONTENTS

					Page
1.	INTR	ODUCTION		• • •	10
2.	CURI	RENT SCOPE OF WORK			14
	2.1	Task 1: In Vitro Primary Antiviral Evaluations			14
	2.2	Task 2: In Vivo Antiviral Evaluations			. 14
	2.3	Task 3: Secondary Evaluations			
3.	EXPE	ERIMENTAL METHODS			. 15
	3.1	Cell Culture			. 15
	3.2	Test Compounds			-
	J.2	3.2.1 Receipt, Cataloging and Storage			
		3.2.2 Determination of Drug Solubility		• • •	. 16
		3.2.3 Compound Preparation for Testing		• • •	. 16
	3.3	In Vitro Antiviral Evaluations: DNA, Exotic RNA and Retro-Viruses			
	3.3	3.3.1 Vaccinia Virus (VV)			
		3.3.2 Adenovirus Type 2 (AD2)			
		3.3.3 Yellow Fever (YF)			
		3.3.4 Japanese Encephalitis Virus (JE)	• • •	• • •	. 25
			• • •	• • •	. 25
		3.3.6 Punta Toro Virus (PT)			
		3.3.7 Sandfly Virus (SF)			
		3.3.8 Hantaan Virus (HTN)			
		3.3.9 Pichinde Virus (PIC)	• • •	• • •	. 28
		3.3.10 Vesicular Stomatitis Virus (VSV)			
		3.3.11 In Vitro Antiviral Screening: Retroviruses			
	3.4	Prescreen Assays (YF, VE, PT)			
	3.5	Antiviral Evaluation In Vivo			. 38
		3.5.1 Pichinde Virus (PIC) in MHA Hamster			
		3.5.2 Venezuelan Equine Encephalomyelitis Virus (VE)			
		3.5.3 Japanese Encephalitis Virus (JE)			
		3.5.4 Vaccinia Virus (VV)			. 39
4.	RESU	JLTS		• • •	. 41
	4.1	In Vitro Antiviral Evaluations: DNA, RNA and Retroviruses Viruses			
		4.1.1 Vaccinia Virus (VV)			
		4.1.2 Adenovirus Type 2 (AD2)			. 73
		4.1.3 Yellow Fever Virus (YF)			. 79
		4.1.4 Japanese Encephalitis Virus (JE)			
		4.1.5 Venezuelan Equine Encephalomyelitis Virus (VE)			. 144
		4.1.6 Punta Toro Virus (PT)			
		4.1.7 Sandfly Virus (SF)			
		4.1.8 Hantaan Virus (HTN)			
		4.1.9 Pichinde Virus (PIC)			
		4.1.10 Vesicular Stomatitis Virus (VSV)			
		4.1.11 In Vitro Screening: Retroviruses			

TABLE OF CONTENTS (CONT'D)

					Page
	4.2	In Vitre	o Prescreen Antiviral Evaluation	 	 294
		4.2.1	Yellow Fever Virus (YF)	 	 296
		4.2.2	Venezuelan Equine Encephalomyelitis Virus (VE)		309
		4.2.3	Punta Toro Virus (PT)		
	4.3	Antivir	al Evaluation In Vivo		
		4.3.1	Pichinde Virus in MHA Hamster		
		4.3.2			
		4.3.3	Japanese Encephalitis Virus (JE)		
		4.3.4	Vaccinia Virus (VV)	 • •	 403
5.	DISC	USSION		 	 406
6.	ACK	NOWLE	OGMENTS	 • •	 408
7.	LITE	RATURE	CITED	 	 409
8.	ABST	RACTS/	PUBLICATIONS	 	 410
9.	LIST	OF PER	SONNEL RECEIVING CONTRACT SUPPORT	 	 411
APPE	NDICE:	S (See V	olume II)		412

LIST OF TABLES

	Page
Table 1	In Vitro Antiviral Screening Models
Table 2	Number of Compounds Received through January 31, 1991
Table 3	AVS Compounds Active Against VV Virus at AI ₉₅ Level
Table 4	AVS Compounds Active Against VV Virus at Also Level
Table 5	Confirmatory Assays for Compounds Active Against Vaccinia Virus 59
Table 6	Confirmatory Assays for Compounds Active Against Adenovirus Type 2 75
Table 7	AVS Compounds Active Against YF Virus at Al ₉₅ Level
Table 8	AVS Compounds Active Against YF Virus at Also Level
Table 9	Confirmatory Assays for Compounds Active Against
- 121	Yellow Fever Virus
Table 10	AVS Compounds Active Against JE Virus at AI ₉₅ Level
Table 11	AVS Compounds Active Against JE Virus at AI ₅₀ Level
Table 12	Confirmatory Assays for Compounds Active Against
	Japanese Encephalitis Virus
Table 13	AVS Compounds Active Against VE Virus at AI ₉₅ Level
Table 14	AVS Compounds Active Against VE Virus at AI ₅₀ Level
Table 15	Confirmatory Assays for Compounds Active Against
	Venezuelan Equine Encephalomyelitis 163
Table 16	AVS Compounds Active Against PT Virus at AI ₉₅ Level
Table 17	AVS Compounds Active Against PT Virus at AI ₅₀ Level
Table 18	Confirmatory Assays for Compounds Active Against Punta Toro 195
Table 19	AVS Compounds Active Against SF Virus at Al ₉₅ Level
Table 20	AVS Compounds Active Against SF Virus at AI ₅₀ Level
Table 21	Confirmatory Assays for Compounds Active Against Sandfly Fever Virus 243
Table 22	Compounds Active Against Hantaan Virus
Table 23	Compounds Active Against Pichinde Virus
Table 24	Confirmatory Assays for Compounds Found Active Against Pichinde Virus
Table 25	Confirmatory Assays for Compounds Active Against
Table 25	Vesicular Stomatitis Virus
Table 26	Compounds Active Against Human Immunodeficiency Virus
Table 27	Confirmatory Assays for Compounds Active Against
m 11 00	Human Immunodeficiency Virus
Table 28	Retesting of AVS-999 Samples
Table 29	Compounds Active Against Feline Leukemia Virus FAIDS Variant
Table 30	Secondary Testing of AVS Compounds Showing Confirmed Activity
T 11 21	Against HIV in ATH8 Cells Assays
Table 31	New Prescreen Drugs that Produced 50% Antiviral Reduction
m 11 20	Against YF Virus
Table 32	New Prescreen Drugs that Produced 50% Antiviral Reduction
m.11 22	Against VE Virus
Table 33	New Prescreen Drugs that Produced 50% Antiviral Reduction Against PT Virus
Table 34	Confirmatory Testing of Compounds Selected From Prescreen Testing 339
Table 35	Compounds Assessed Against Pichinde Virus In Vivo
Table 36	Antiviral Efficacy of AVS-206
Table 37	Results of Retesting of AVS-206
Table 38	Antiviral Efficacy of AVS-206 and Ribavirin (AVS-01)
Table 39	Antiviral Efficacy of AVS-1046

LIST OF TABLES (CONT'D)

	<u>Page</u>
Table 40	Compounds Received for Testing Against Venezuelan Equine Encephalomyelitis Virus
Table 41	Cumulative Results of Intracranial Challenge with Venezuelan
	Equine Encephalomyelitis Virus in Mice
Table 42	Mortality in VEE Challenged Mice Receiving a Single Dose of 3N-3DU 393
Table 43	Compounds Received for Testing Against Japanese Encephalitis Virus In Vivo 396
Table 44	Antiviral Evaluation of AVS-360 (JE Virus)
Table 45	In Vivo Antiviral Efficacy of AVS-360 and AVS-2563 Against JE Virus 398
Table 46	Antiviral Efficacy of AVS-361, 2811 and 2812 Against JE Virus 399
Table 47	Antiviral Efficacy of Selected AVS Compounds Against JE Virus 400
Table 48	In Vivo Testing of AVS-2979 Against JE Virus Challenge
Table 49	Results of Testing of AVS-5587 Against Japanese Encephalitis
	Virus Challenge
Table 50	Compounds Received for Testing Against In Vivo Vaccinia Virus
Table 51	Activity of Selected AVS Compounds vs Vaccinia Virus (Tailpox Counts) 405

LIST OF FIGURES

		Page
Figure 1	Primary-Screen Data Sheet	20
Figure 2	Prescreen Data Sheet	
Figure 3	Total Number of In Vitro Antiviral Assays Performed per Year	
1 -841-0	(November 16, 1985 - January 31, 1991)	43
Figure 4	Number of Compounds Tested In Vitro Per Year During the Contract Period	
Figure 5	In Vitro Primary Screen: Number of Compounds Found Active	
rigute 3	Against Vaccinia Virus During the Contract Period	45
F: C A		
Figure 6-A	Ara-A - vs - Vaccinia Virus; Values for 89 Positive Control Assays	. 47
Figure 6-B	Selenazofurin - vs - Vaccinia Virus; Values for 83 Positive Control	40
	Assays	49
Figure 7	Variation of the Maximum Antiviral Effect of Vaccinia Virus - vs -	
	Ara-A/Selenazofurin	50
Figure 8	Variation of the Cell (Load) Controls; Vaccinia Virus - vs -	
	Ara-A/Selenazofurin	51
Figure 9	Variation of the Virus (Load) Controls; Vaccinia Virus - vs -	
	Ara-A/Selenazofurin	51
Figure 10	Variation of the Test Differential; Vaccinia Virus - vs -	
	Ara-A/Selenazofurin	51
Figure 11	In Vitro Primary Screen: Number of Compounds Found Active	
	Against Adenovirus During the Contract Period	73
Figure 12	In Vitro Primary Screen: Number of Compounds Found Active	
1 .5410 12	Against Yellow Fever Virus During the Contract Period	79
Figure 13-A		, ,,
I iguic 15-A		81
Figure 12 D	Assays	01
Figure 13-B		06
F: 14 A	Compound Tests	86
Figure 14-A		00
	Selenazofurin	82
Figure 14-B	Variation of Maximum Antiviral Effect; Yellow Fever Virus - vs -	
	2-Thio-6-Azauridine	
Figure 15-A	Variation of Cell (Load) Controls; Yellow Fever Virus - vs - Selenazofurin	. 84
Figure 15-B	Variation of Cell (Load) Controls; Yellow Fever Virus - vs -	
	2-Thio-6-Azauridine	89
Figure 16-A	Variation of Virus (Load) Controls; Yellow Fever Virus - vs - Selenazofurin	. 84
Figure 16-B	Variation of Virus (Load) Controls; Yellow Fever Virus - vs -	
	2-Thio-6-Azauridine	89
Figure 17-A	Variation of Test Differential; Yellow Fever Virus - vs - Selenazofurin	
Figure 17-B		
Figure 18	In Vitro Primary Screen: Number of Compounds Found Active	. 02
1.5410 10	Against Japanese Encephalitis Virus During the Contract Period	115
Figure 19-A		113
rigule 19-A		117
Figure 10 D	Control Compound Tests	117
rigure 19-B	2-Thio-6-Azauridine - vs - Japanese Encephalitis Virus; Values for 46	
E: 00 :	Positive Control Compound Tests	122
Figure 20-A	Variation of Maximum Antiviral Effect; Japanese Encephalitis Virus - vs -	
	Selenazofurin	118
Figure 20-B	Variation of Maximum Antiviral Effect: Japanese Encephalitis Virus - vs -	
	2-Thio-6-Azauridine	123

LIST OF FIGURES (CONT'D)

		Page
Figure 21-A	Variation of Cell (Load) Controls; Japanese Encephalitis Virus - vs -	120
Figure 21-B	Selenazofurin	120
Figure 22-A	Azauridine	
Figure 22-B	Selenazofurin	120
Figure 23-A	2-Thio-6-Azauridine	125
Figure 23-B	Selenazofurin	120
Figure 24	2-Thio-6-Azauridine	125
	Against Venezuelan Equine Encephalomyelitis Virus During the Contract Period Selenazofurin - vs - Venezuelan Equine Encephalomyelitis Virus;	144
	Values for 220 Positive Control Compound Tests	146
	2-Thio-6-Azauridine - vs - Venezuelan Equine Encephalomyelitis Virus; Values for 42 Positive Control Compound Tests	151
	Variation of Maximum Antiviral Effect; Venezuelan Equine Encephalomyelitis Virus - vs - Selenazofurin	147
	Variation of Maximum Antiviral Effect; Venezuelan Equine Encephalomyelitis Virus - vs - 2-Thio-6-Azauridine	152
Figure 27-A	Variation of Cell (Load) Controls; Venezuelan Equine Encephalomyelitis Virus - vs - Selenazofurin	149
Figure 27-B	Variation of Cell (Load) Controls; Venezuelan Equine Encephalomyelitis Virus - vs - 2-Thio-6-Azauridine	154
Figure 28-A	Variation of Virus (Load) Controls; Venezuelan Equine Encephalomyelitis Virus - vs - Selenazofurin	149
Figure 28-B	Variation of Virus (Load) Controls; Venezuelan Equine Encephalomyelitis Virus - vs - 2-Thio-6-Azauridine	
Figure 29-A	Variation of Test Differential; Venezuelan Equine Encephalomyelitis Virus - vs - Selenazofurin	
Figure 29-B	Variation of Test Differential; Venezuelan Equine Encephalomyelitis Virus - vs - 2-Thio-6-Azauridine	
Figure 30	In Vitro Primary Screen: Number of Compounds Found Active Against Punta Toro Virus During the Contract Period	
Figure 31-A	Ribavirin - vs - Punta Toro Virus; Values for 242 Positive Control	
Figure 31-B	Compound Tests	
	Variation of Maximum Antiviral Effect; Punta Toro Virus - vs - 2-Thio-6-Azauridine	
	Variation of Cell (Load) Controls; Punta Toro Virus - vs - Ribavirin	
	Variation of Cell (Load) Controls; Punta Toro Virus - vs - 2-Thio-6-Azauridine	
	Variation of Virus (Load) Controls; Punta Toro Virus - vs - Ribavirin	
Figure 34-B	Variation of Virus (Load) Controls; Punta Toro Virus - vs - 2-Thio-6-Azauridine	
Figure 35-A	Variation of Test Differential; Punta Toro Virus - vs - Ribavirin	178

LIST OF FIGURES (CONT'D)

Pag	ze
Wasietian of Tax Differentials Dante Tons Views are 2 This 6 Amountains 19	22
In Vitro Primary Screen: Number of Compounds Found Active Against	
2-Thio-6-Azauridine - vs - Sandfly Virus; Values for 42 Positive	
In Vitro Primary Screen: Number of Compounds Found Active Against	
	8
	16
	6
	8
)5
	16
	8
· · · · · · · · · · · · · · · · · · ·)3
	9
)4
)1
)6
)1
)6
)1
(Prescreen Protocol)	16
	Variation of Test Differential; Punta Toro Virus - vs - 2-Thio-6-Azauridine In Vitro Primary Screen: Number of Compounds Found Active Against Sandfly Fever Virus During the Contract Period Ribavirin - vs - Sandfly Virus; Values for 226 Positive Control Compound Tests 227-Thio-6-Azauridine - vs - Sandfly Virus; Values for 42 Positive Control Compound Tests 228 Variation of Maximum Antiviral Effect; Sandfly - vs - Ribavirin 229 Variation of Maximum Antiviral Effect; Sandfly - vs - Ribavirin 220 Variation of Cell (Load) Controls; Sandfly - vs - Ribavirin 221 Variation of Cell (Load) Controls; Sandfly - vs - Ribavirin 222 Variation of Virus (Load) Controls; Sandfly - vs - Ribavirin 223 Variation of Virus (Load) Controls; Sandfly - vs - Ribavirin 224 Variation of Test Differential; Sandfly - vs - Ribavirin 225 Variation of Test Differential; Sandfly - vs - Ribavirin 226 Variation of Test Differential; Sandfly - vs - Ribavirin 227 Variation of Test Differential; Sandfly - vs - Ribavirin 228 In Vitro Primary Screen: Number of Compounds Found Active Against 239 In Vitro Primary Screen: Number of Compounds Found Active Against 240 Variation of Test Differential; Sandfly - vs - Z-Thio-6-Azauridine 251 In Vitro Primary Screen: Number of Compounds Found Active Against 252 Variation of Test Differential; Sandfly - vs - Z-Thio-6-Azauridine 253 Variation of Test Differential; Sandfly - vs - Z-Thio-6-Azauridine 264 Nativity of Nativity of Nativity Activity of Nativity Of Na

LIST OF FIGURES (CONT'D)

		Page
Figure 56	In Vitro Prescreen: Number of Compounds Found Active Against	
	Venezuelan Equine Encephalomyelitis Virus During the Contract Period	309
Figure 57-A	Selenazofurin - vs - Venezuelan Equine Encephalomyelitis Virus (Prescreen Protocol)	311
Figure 57-B		
	(Prescreen Protocol)	316
Figure 58-A	Variation of Maximum Antiviral Effect; Venezuelan Equine	
T: 60 D	Encephalomyelitis Virus (Prescreen Protocol)	312
Figure 58-B	Variation of Maximum Antiviral Effect; Venezuelan Equine Encephalomyelitis Virus (Prescreen Protocol)	217
Figure 59-A	Variation of Cell (Load) Controls; Venezuelan Equine	517
8	Encephalomyelitis Virus - vs - Selenazofurin (Prescreen Protocol)	314
Figure 59-B	Variation of Cell (Load) Controls; Venezuelan Equine Encephalomyelitis	
	Virus - vs - 2-Thio-6- Azauridine (Prescreen Protocol)	319
Figure 60-A	Variation of Virus (Load) Controls; Venezuelan Equine Encephalomyelitis	214
Figure 60 B	Virus - vs - Selenazofurin (Prescreen Protocol)	314
rigule 00-D	Virus - vs - 2-Thio-6-Azauridine (Prescreen Protocol)	319
Figure 61-A	Variation of Test Differential; Venezuelan Equine Encephalomyelitis	
9	Virus - vs - Selenazofurin (Prescreen Protocol)	314
Figure 61-B	Variation of Test Differential; Venezuelan Equine Encephalomyelitis	
	Virus - vs - 2-Thio-6-Azauridine (Prescreen Protocol)	319
Figure 62	In Vitro Prescreen: Number of Compounds Found Active Against	22.1
Figure 63-A	Punta Toro Virus During this Contract Period	321
rigule 05-A	Control Compounds Assays	323
Figure 63-B	2-Thio-6-Azauridine - vs - Punta Toro Virus; Values for 60 Positive Control	
	Compounds Assays	328
Figure 64-A	Variation of Maximum Antiviral Effect; Punta Toro Virus - vs - Ribavirin	
E' (4 P	(Prescreen Protocol)	324
Figure 04-B	Variation of Maximum Antiviral Effect; Punta Toro Virus - vs - 2-Thio-6-Azauridine (Prescreen Protocol)	320
Figure 65-A	Variation of Cell (Load) Controls; Punta Toro Virus - vs -	329
1 18410 00 11	Ribavirin (Prescreen Protocol)	326
Figure 65-B	Variation of Cell (Load) Controls; Punta Toro Virus - vs -	
	2-Thio-6-Azauridine (Prescreen Protocol)	331
Figure 66-A	Variation of Virus (Load) Controls; Punta Toro Virus - vs - Ribavirin	224
Figure 66 D	(Prescreen Protocol)	326
rigure 00-b	Variation of Virus (Load) Controls; Punta Toro Virus - vs - 2-Thio-6-Azauridine (Prescreen Protocol)	331
Figure 67-A	Variation of Test Differential; Punta Toro Virus - vs - Ribavirin	551
	(Prescreen Protocol)	326
Figure 67-B	Variation of Test Differential; Punta Toro Virus - vs - 2-Thio-6-Azauridine	
	(Prescreen Protocol)	331

1. INTRODUCTION

This is the Final Progress Report on SRI Project No. 5975, Contract No. DAMD17-86-C-6013. It covers the progress of the research program during the report period from November 16, 1985 to January 31, 1991.

The goal of this program was to implement testing systems in which to evaluate the efficacy of candidate antiviral compounds against a spectrum of viruses of military importance. The program consists of three major task areas: a) primary testing of chemical compounds and natural products for antiviral efficacy in vitro, b) primary testing of chemical compounds and immunopotentiators for antiviral efficacy in vivo, and c) secondary evaluation of compounds found active in the primary in vitro and in vivo screens.

One of the primary missions of the U.S. Army Medical Research and Development Command is to perform studies on the pathogenesis, diagnosis, epidemiology, prophylaxis, and treatment of infectious diseases of military importance. The Army's infectious disease research program, conducted by the U.S. Army Medical Research Institute of Infectious Diseases (USAMRIID) at Fort Detrick, is primarily concerned with medical defense against (a) naturally-occurring infectious diseases that could seriously interrupt U.S. military operations such as troop mobilization and deployment and (b) the threat of infectious diseases or toxic effects caused by the potential field use of biological warfare (BW) agents, either conventional BW agents or altered agents, by an unfriendly force.

The U.S. Army has a recognized need for new chemical compounds that will be useful as prophylactic or therapeutic antiviral drugs to treat U.S. military personnel who are at risk of exposure to, or who might become infected with, naturally-occurring viruses or altered viruses for which there exists no effective protection or therapy at the present time. The development of selective antiviral drugs for use in the successful treatment of infections with certain exotic RNA viruses (togaviruses, bunyaviruses, arenaviruses, rhabdoviruses, and other, unclassified RNA viruses) is of particular importance to the Army because there are no other research efforts being conducted, either by the government or by the private sector, which are directed toward the control of these virus diseases of military relevance.

In 1973, USAMRIID initiated a research and development program to identify and to pursue new compounds with activity against these exotic RNA viruses. Approximately 1500 compounds have already been screened in vitro for selective antiviral effects against these target viruses and a number of the compounds which were found active in cell culture have been evaluated for antiviral efficacy in vivo. Several of these compounds (e.g., Ribavirin, Selenazole, and Pyrazofurin) have been extensively tested for efficacy against lethal RNA virus infections in various animal model systems at USAMRIID. To date, the most promising antiviral drug with demonstrated, broad-spectrum activity against these viruses of military importance, both in vitro and in vivo, appears to be Ribavirin, its prodrug derivatives, and its carboxamidine derivative (AVS-206). Ribavirin has been evaluated in humans infected with Sandfly Fever (SF) Virus, Lassa Fever Virus, and Korean Hemorrhagic Fever (KHF) Virus and has demonstrated marked clinical efficacy against these particular virus infections. This drug will be further developed for general use in military personnel. There is a real need, however, for more potent and more selective antiviral drugs to combat these virus diseases which represent serious threats throughout the world.

Troops in the field are threatened not only by infectious diseases of natural origin, but also by the possibility of BW attack. The commercial development of antiviral drugs for the treatment of the

more common respiratory virus, enteric virus, and herpesvirus infections may not solve the problems which are unique to the Armed Forces. These potential problems of encountering exotic viruses and BW agents will not be sufficiently addressed by depending solely on the possibility that antiviral drugs originally developed for the treatment of acute respiratory diseases, enterovirus infections, and herpesvirus infections might also be useful in the treatment of these virus diseases of military relevance. A more direct approach, and one which is clearly indicated, is to focus on antiviral drug development efforts designed to attack these particular virus disease threats that are unique to the Armed Forces.

The U.S. Army Medical Research Institute of Infectious Diseases has, for a number of years, been involved in conducting a unique and ambitious antiviral drug research and development program, primarily directed toward the chemical control of exotic RNA virus infections of military importance. Potential antiviral agents have been synthesized and evaluated against a number of target viruses both in vitro and in vivo. Program emphasis is currently on the development of antivirals for use in the treatment of infections with alphaviruses, flaviviruses, bunyaviruses, arenaviruses, and other viruses which are capable of eroding combat strength in troops deployed in overseas areas. In addition, current efforts are also being directed toward the development of antivirals for use in the treatment of AIDS through an Inter-Agency Agreement with the National Institutes of Health (NIH).

Members of the *Togaviridae* and *Flaviviridae* families (alphaviruses and flaviviruses) are capable of producing serious hemorrhagic or encephalitic diseases in humans. Infections with alphaviruses [Eastern equine encephalomyelitis (EEE), Western equine encephalitis (WEE), and Venezuelan equine encephalitis (VEE)] have occurred in epidemic proportions in the Americas. Chikungunya and O'nyongnyong viruses also continue to cause epidemic disease on the African continent. The flaviviruses include several members which cause significant disease in humans. Dengue viruses types 1-4 are prevalent causes of acute illness in the tropics and subtropics of the world. Available vaccines are inadequate to control these infections effectively. Other members such as St. Louis encephalitis virus, Japanese B Encephalitis (JE) virus, and West Nile encephalitis virus cause mild to severe encephalitis diseases in humans. The tick-borne encephalitis virus group, represented by Russian Spring-Summer Encephalitis Virus, has caused widespread encephalitic disease in the U.S.S.R. and Northern Europe with high mortality rates. Yellow Fever (YF), in either the urban or jungle form, continues to be a threat, although the use of the 17-D vaccine is quite effective as a prophylactic measure against this disease. The Army's program is interested in controlling infections caused by the dengue viruses, Japanese encephalitis virus, Russian Spring-Summer encephalitis virus, Yellow Fever virus, and West Nile encephalitis virus.

A number of bunyaviruses have caused epidemic disease in many areas of the world: Rift Valley Fever virus was responsible for a major epizootic in Egypt in 1977-79 with considerable losses in domestic animals (sheep and cattle) and significant mortality among those humans infected with the virus. Sandfly fever virus has also been recognized as an important cause of epidemics in the Mediterranean area, in the Middle East, and in Central Europe. Oropouche, La Crosse, and California encephalitis viruses have all caused significant disease in the Americas. Oropouche virus, for example, has been associated with a number of large human epidemics in Brazil over the past twenty years. Hantaan virus, the causative agent of Korean hemorrhagic fever, causes appreciable mortality and is widely distributed in Asia. It has only recently been shown to belong to the Bunyaviridae family. Of the Bunyaviridae family, the USAMRIID program has initiated studies with Sandfly fever virus, Rift Valley fever virus, Korean hemorrhagic fever virus, and Punta Toro virus.

Of the Arenaviridae family, current interest includes Lassa fever virus, an agent which causes significant lethal disease among infected humans in Africa, especially in Sierra Leone. Other arenaviruses

under current investigation include Junin and Machupo viruses, the causative agents of Argentine hemorrhagic fever and Bolivian hemorrhagic fever, respectively. These agents are found endemic in wide areas of South America. Pichinde (PIC) virus has been used in antiviral studies as a representative of this important family of viruses. Vesicular stomatitis virus is currently employed as a representative of the *Rhabdoviridae* family. Vaccinia Virus is currently employed as a representation of the DNA Virus (Poxviridae). This agent poses a threat to the military personnel as well as the general population because of the lack of antibody protection since no World Health Vaccination Program are now required. VV Virus is also commonly used as a carrier virus for genetic engineering technology, therefore posing an ever-present threat to a laboratory modification of any genetically engineered virulent agent.

Other viruses which cause sporadic but severe hemorrhagic fever in Africa are Marburg and Ebola viruses. These two closely-related agents have been placed in a new family (Filoviridae).

The above viruses are those which might be encountered in exotic troop locations and against which troops would not be expected to have any pre-existing immunity. With few exceptions, specific vaccines do not exist for these agents and some of the agents encountered may be poorly classified or may even be unclassified viruses which have not been seen previously. The antiviral chemoprophylaxis/chemotherapy approach may be the best modality to defend against this threat at the present time.

Another recurring problem with naturally-occurring virus infections exists in military boot camps where new recruits are assembled. These troops often develop infections with adenoviruses, influenza viruses, and parainfluenza viruses, sometimes in epidemic proportions. Infections with the adenoviruses have been a distinct military problem for years and multivalent vaccines have been prepared for use in new recruits. Nevertheless, an effective antiviral drug for treatment of adenovirus infections would be quite useful and therefore this virus group is also a target for antiviral chemotherapy in the Army's program.

A number of naturally-occurring viruses could be developed by an adversary into potent biological weapons for use in the field against U.S. forces. Many of the exotic RNA viruses are also potential BW agents, since their dissemination in an area where they are already indigenous could be employed as a means of disguising the source of the infection. In addition, other agents such as smallpox virus could be used very effectively against a susceptible civilian population prior to and during military operations to disrupt logistics and support activities and to create panic and chaos. The threat of BW attack poses some of the same problems as those to be addressed in the defense against naturally-occurring virus infections. Again, a broad-spectrum antiviral drug with selective activity against the RNA viruses could be the only real line of defense against such attach with those particular type of agents. The vaccine approach will only be effective in affording protection against a very limited group of these agents which number in the hundreds of different antigenic types.

With the advances made in the field of molecular genetics, it is now technically possible to genetically engineer altered viruses with enhanced virulence, communicability, drug resistance, and overall threat potential. An example of such misuse of advanced biotechnology would be the insertion of genes for highly toxic peptides such as snake venom toxins, potent bacterial toxins, or other low molecular weight toxins of military importance into the genome of a highly communicable virus such as influenza. The feasibility of inserting foreign genes into vaccinia virus and obtaining expression of those genes in the host cell has already been demonstrated. Recombinant DNA technology has made it possible to insert and express heterologous genes in a variety of different viruses. Effective defense against

possible altered viruses used as BW agents may well depend upon the development of antiviral drugs active against the vector viruses.

The Department of Antiviral Studies at USAMRIID is responsible for the acquisition, identification, and development of potential new antiviral drugs which are effective against viruses of military relevance and which might be useful in the treatment of AIDS. The program is therefore, broad-based and involves the synthesis, primary and secondary testing, and further characterization of novel antiviral compounds with regard to their possible biochemical mechanisms of action, pharmacokinetics, metabolism, optimal formulation, optimal combination with other drugs, and safety in animal model systems. The Department also directs studies in support of IND applications to the FDA for clinical testing of active antiviral agents for use in man.

Since the establishment of its antiviral testing program, the Department of Antiviral Studies, USAMRIID, has evaluated approximately 3,000 compounds in its primary in vitro screen. Fewer compounds have been evaluated in vivo. Selenazole was reported to have broad-spectrum activity against the exotic RNA viruses in vitro and is significantly more potent than Ribavirin against the togaviruses (VEE, YF, JE), bunyaviruses (RVF, SF, KHF), and arenaviruses (PIC) in vitro. The activity of this compound against YF virus is most impressive with an ED₅₀ of 0.005 mg/ml in cell culture. The evaluation of Selenazole for therapeutic antiviral efficacy in vivo, however, yielded disappointing results and pharmacological problems may be responsible for the lack of efficacy in animal model systems. Ribavirin, on the other hand, has been shown in laboratory animal models to have significant antiviral efficacy against the bunyaviruses (RFV, Punta Toro and KHF) and the arenaviruses (PIC, Junin, Machupo, and Lassa Fever). Clinical trials with Ribavirin in patients with Lassa Fever virus or Hantavirus infections have yielded good results.

Ribavirin has also been shown to be effective in the treatment of Sandfly fever virus in human volunteers. Good progress has been made toward developing this particular antiviral drug for general clinical use by the Army, but new agents with higher potency and selectivity against the exotic RNA viruses will hopefully be identified in the expanded USAMRIID antiviral program. Because of the lag time in diagnosing viral disease, treatment with broad-spectrum antiviral agents offers the best hope to successfully defend against both naturally-occurring disease and against possible BW agents in the field. It is unlikely, however, that a single drug will be found that is effective against all of these exotic RNA virus infections, so additional antiviral agents must be developed. There is also a need to explore the efficacy of immunopotentiators, biological response modifiers, interferons, combination chemotherapy, and new approaches to drug delivery to enhance the antiviral efficacy of these agents.

The basic contract at Southern Research Institute was established to enable USAMRIID to evaluate approximately 1,500 compounds per year for efficacy against 10 different target viruses in a primary in vitro screen, 96 compounds per year for in vivo efficacy against a representative togavirus and arenavirus in appropriate animal model systems, and approximately 5 compounds per year in detailed in vivo studies. The basic contract also includes secondary testing studies with candidate antiviral agents that demonstrate promising activity both in vitro and in vivo. In July, 1986, the HIV virus was added as an additional virus to be tested under the in vitro primary screen. During 1989, at the request of Ms. C. Susan Tiffany (Contract Specialist, Ft. Detrick, Frederick, Maryland) the Adenovirus (AD2) and the Vesicular Stomatitis Virus (VSV) were deleted from the primary screen. Also, all in vivo work was discontinued. During 1989 - 1990, antiviral prescreen testing (in vitro) of plant extracts against Punta Toro, Yellow Fever Virus and Venezuelan Equine Encephalitis was added to this screening program at the rate of approximately 3000 extracts per year. This report summarizes our progress in implementing the research program and includes summaries of antiviral test data collected from November 16, 1985, through January 31, 1991.

2. CURRENT SCOPE OF WORK

This section describes the research objectives and the scope of work for each of the main tasks being performed by Southern Research Institute (SRI) on this contract during this reporting period. These tasks are: (a) Primary testing of compounds and plant extracts for antiviral efficacy in vitro, (b) Primary testing of chemical compounds and immunopotentiators for antiviral efficacy in vivo, (c) Secondary evaluation of compounds found active in the primary in vitro and in vivo screens.

2.1 Task 1: In Vitro Antiviral Evaluations

SRI conducted the primary screening of chemical compounds which were furnished by the Department of Antiviral Studies, USAMRIID, through its repository contractor (Biological Research Faculty and Facility, Inc., [BRFF, Inc.] Ijamsville, MD) for antiviral efficacy in cell culture against representative viruses from the Togaviridae, Flaviviridae, Bunyaviridae, Arenaviridae, Rhabdoviridae, Poxviridae, Adenoviridae and Retroviridae families. The test viruses consist of the following: (1) Vaccinia (VV) Virus, (2) Adenovirus Type 2 (AD2), (3) Yellow Fever (YF) Virus, (4) Japanese Encephalitis (JE) Virus, (5) Venezuelan Equine Encephalomyelitis (VE) Virus, (6) Punta Toro (PT) Virus, (7) Sandfly (SF) Virus, (8) Hantaan (HTN) Virus, (9) Pichinde (PIC) Virus, (10) Vesicular Stomatitis (VSV) Virus, (11) Human Immunodeficiency (HIV) Virus. SRI was scheduled to evaluate approximately 1500 compounds per year against each of these eleven viruses in vitro, using CPE-inhibition assays, to determine the 50% minimal inhibitory concentration (MIC₅₀), or median inhibition dose (ID₅₀) of active compounds, respectively. Determinations was also made of the cytotoxicity of each candidate compound for the host cells, expressed in terms of the minimum ($\sim 25\%$) cytotoxic concentration of the compound. We calculated an in vitro selectivity index for each active compound against each susceptible test virus. Compounds that showed antiviral activity in the initial CPE-inhibition were retested in confirmatory CPEinhibition assays.

Our program was changed during the latter part of 1989 to implement the Antiviral Prescreen Assay to evaluate plant extracts on a large scale against PT, YF and VE viruses. We were scheduled to test approximately 2000 - 3000 extracts per year using the prescreen assay protocol. A prescreen procedure was developed which uses MTT and evaluates five compounds (at four-log₁₀-dose levels) per virus per 96-well plate. Compounds deemed active from the prescreen program were tested further in the primary screening program against the six Exotic Viruses listed above. Screening data was reported to BRFF, Inc. and to USAMRIID essentially as it was obtained in hardcopy form and/or on floppy diskettes.

2.2 Task 2: In Vivo Antiviral Evaluations

SRI conducted the primary testing of chemical compounds and immunopotentiators, being furnished by the Department of Antiviral Studies, USAMRIID, for antiviral efficacy in rodent models and conducted a preliminary assessment of acute toxicity with each material submitted. SRI evaluated the compounds and immunopotentiators against representative viruses from the Togaviridae and Arenaviridae families in rodents. These challenge viruses consist of the following agents: (1) Venezuelan Equine Encephalomyelitis Virus in the mouse, and (2) Pichinde Virus in the hamster. It was later decided to expand the *in vivo* evaluations to include representative viruses from the Flaviviridae, Poxviridae, and Retroviridae families. This expanded our list of challenge viruses to include the following agents: (3) Japanese Encephalitis Virus and (4) Vaccinia Virus. Compounds are evaluated at the primary level against each of these viruses using parenteral routes of administration. Antiviral efficacy was expressed in terms of the observed increase in the number of survivors, or in the mean survival time, in the treated group compared with that of the control untreated group. An *in vivo* virus rating was determined.

2.3 Task 3: Secondary Evaluations

SRI also conducted the secondary testing and further evaluation of compounds found active in the primary antiviral screen (Confirmatory Assays). Secondary testing includes an estimation of the in vivo therapeutic index of the candidate compound, the optimal dosage regimen, the optimal route of administration, the optimal treatment schedule, schedule dependency, the duration of antiviral effect, a determination of the utility of the candidate compound in combination with other drugs, and the influence of formulation on activity. SRI is expecting to evaluate approximately 5 compounds per year in extensive secondary in vivo testing to include determinations of dose-response relationships and antiviral efficacy. Data was reported to USAMRIID essentially as it was obtained.

Secondary in vitro evaluations of compounds active against Human Immunodeficiency Virus in the primary screen were also performed. These compounds were examined for their ability to inhibit HIV-induced p24 gag protein expression in permissive target cells by means of an indirect immunofluorescence assay, and for their ability to inhibit the production of HIV-induced reverse transcriptase activity in infected target cells (see section 4 of the Second Annual Report dated December 12, 1987). Active compounds are also being tested for possible cytotoxic effects in various immune function assays. In addition, active compounds were tested in vitro for efficacy against related murine, feline, and simian retroviruses.

3. EXPERIMENTAL METHODS

3.1 Cell Culture

A centralized cell culture facility for provision of high-quality cell culture for all of the virus screening laboratories on this project is located at SRI home-site. This centralized cell culture facility consists of two laboratories (one laboratory is approximately 225 sq. ft. and the other approximately 250 sq. ft.). Each laboratory has a laminar flow hood and each laboratory is fully equipped for the maintenance and propagation of cell cultures. Each laminar flow hood has been inspected and certified by the University of Alabama Occupational Health and Safety Department and these inspections and certifications will be done on a yearly basis. One laboratory has been in operation since February 1988 and the other laboratory was completed and came on line in June, 1988.

The centralized cell culture facility is currently propagated and maintained the following cell culture lines for this project: Vero (ATCC) and LLC-MK2.

Vero (ATCC) cells are used for seeding 96-well plates for the RNA virus assays and delivered to the BL-3 facility on the following schedules. Cell culture plates for the other virus laboratories are supplied as requested. Vero cells in T75 or T150 cell culture flasks for virus production are supplied as requested. In addition, Vero cells seeded in 96-well plates are supplied as requested for special studies, and developmental procedures.

The Centralized Cell Culture Facility supplied the media and solutions to the BL-3 facility as was needed.

For quality control, cell culture lines are routinely monitored for any change in their growth parameters. The cells seeded in 96-well plates are microscopically inspected before delivery to the laboratories to ensure proper cell distribution in the wells and cell integrity.

Sterility cultures are performed on all media and solutions used in the cell culture laboratory. At the time each reagent is made, 0.5 ml from each bottle or flask of reagent is added to an individual tube of thioglycollate medium and Sabouraud medium and the sterility culture tubes are incubated at 37°C for 48 hours before the solutions are released for use. The culture tubes are held 14 days before being discarded as negative.

At approximately 3-month intervals, samples from cell culture lines that are maintained in the Centralized Cell Culture Facility are sent to the ATCC for mycoplasma testing. These samples are monitored by use of the bisbenzamide DNA-fluorochrome stain and also by cultivating in mycoplasma broth and agar media under aerobic and anaerobic culture conditions.

Cell culture stocks are stored in liquid nitrogen.

3.2 Test Compounds

3.2.1 Receipt, Cataloging, and Storage

The drug samples submitted for testing were shipped to the centralized drug preparation laboratory by Biological Research Faculty and Facility, Inc. (BRFF). The drugs were checked against the enclosed shipping list, and stored in numerical order in the drug repository facility under the appropriate conditions, according to the information supplied by BRFF, Inc.

Requests for drug preparation are required to be delivered to the centralized drug preparation laboratory five working days prior to (Drug Request Form) the testing date. The drugs are solubilized and delivered to the designated laboratory on the day of testing.

3.2.2 Determination of Drug Solubility

If no drug solubility or stability information is provided by the supplier, the following procedure is used to determine drug solubility:

Weigh a 1 mg sample into a homogenizer vessel and add 1 ml of H_2O , which is the first solvent on the priority list. If the drug is not immediately soluble in H_2O , heat in a H_2O bath to $40^{\circ}C$. If the drug is not in solution after heating, homogenize with a hand homogenizer. Repeat the procedure with a freshly weighed 1 mg sample for each solvent in order of priority, until a suitable solvent is found. The priority list of solvents is as follows:

	Solvent	Volume (ml)		Solvent	Volume (ml)
1)	H ₂ O	1	4)	DMSO	0.1
2)	MeOH	0.1	5)	Acetone	0.1
3)	EtOH	0.1			

If the drug is insoluble in all of the above listed solvents, it is to be tested as a suspension in cell culture assay medium with the aid of a hand homogenizer or a vortex mixer. The final concentration of solvent in the starting drug concentration should not exceed 1% (preferably <1%).

3.2.3 Compound Preparation for Testing

The following procedure is employed for the drug preparation:

1 Drugs for Primary Testing

All drugs are weighed in specified amounts, and the pre-determined solvent is added as required for the starting drug concentration.

Polystyrene snap-cap tubes are used for the weighed samples except for acetone-soluble drugs, then polypropylene tubes must be used. Polypropylene cryotubes are also used.

2. Pre-screen Drugs (Plant Extracts)

Plant extracts are received in the repository preweighed in amounts of 200 mg (± 10%) in 2 ml cryovials with an o-ring seal.

On Day 0 (24 hours prior to testing) 200 μ l of a specified solvent is added to each sample. Each sample is homogenized and placed in an ultrasonic bath for 10 - 12 minutes. The samples are extracted for 18 hours at 22 - 24° C.

On Day 1 (day of testing), 800 μ l of sterile-deionized water is added to each sample and thoroughly mixed. If a drug is insoluble, it is further homogenized with a tissue tearer until a homogenous suspension is obtained. An additional 1 ml of sterile-deionized water is added to each sample, bringing the total volume in each cryovial to 2 ml.

3. NCI Compounds for HIV Testing

Compounds supplied to us from the NCI storage contractor for retrovirus screening were received and prepared for testing in a separate drug preparation laboratory at SRI under the direction of Ms. Patricia Holum.

Tasks, such as acknowledgement of drug receipt, internal data-base management of all drugs, initialization of the *In-vitro* Screening System (IVSS), drug evaluation and data reporting were accomplished with an integrated computer system consisting of local and foreign hardware and software.

Local hardware consisted of 4 micro-computers, 5 Ampex terminals, 2 bar-code printers, 2 dot-matrix printers, 1 laser printer, and 2 96-well plate readers. Local software included public domain communications programs as well as commercial programs for relational data base management and word-processing.

Foreign hardware included two NIH main-frame computers located in Bethesda, Maryland, a DEC 10 and an IBM-370. Access and use of these computers was under the auspices of the Division of Computer Research and Technology (DCRT). Software, the IVSS (*In vitro* screening system), was provided by Value Systems Engineering (VSE) under contract to DCRT.

Initialization of the IVSS was through the AVAIL module of that program on the IBM-370 after drug solubilizations had been performed. Solubility determinations were made using vehicles in the following order; distilled water, DMSO, methyl alcohol, ethyl alcohol, or any other vehicle requested by DTP. When a vehicle was found in which the test agent was per ml. Using the vehicle found in the solubility determination, the test agent was prepared at the highest soluble concentration (maximum 100 mg/ml), labeled with a bar-code label produced in the AVAIL module of the IVSS, and stored at -20°C until delivered to the HIV screening laboratory.

3.3 In Vitro Antiviral Screening: DNA, Exotic RNA Viruses and Retroviruses

The viruses and host cell lines used in our *in vitro* assays are listed in Table 1. The antiviral activity of a compound was defined as a measure of its ability to inhibit the cytopathogenic effect (CPE) of the virus on its host cell. During the first three years of the contract, compounds were evaluated in a standard CPE-inhibition assay (virus rating). The last two years of the program, we moved to a MTT based antiviral assay format. Positive control drugs (Table 1) were included in each antiviral assay to validate the test conditions used in the antiviral assays.

The MTT Assay System, measures the degree of cell viability (and therefore CPE and drug cytotoxicity) as determined by MTT uptake. This procedure is based upon the reduction of the tetrazolium salt, 3-(4,5-dimethyl-thiazol-2-yl)-2,5-diphenyl tetrazolium bromide (MTT) by mitochondrial enzymes of viable host cells to MTT formazan. The blue color of the MTT formazan is measured spectrophotometrically. It was felt that this new system was more efficient and suitable for a large-scale screening program. Plate layout and data printout are included in Figure 1.

The details of the assay procedures for each virus are described on the following pages.

Virus	Strain	Classification	Cell Line P	Positive Control Compound
Vaccinia Virus	Lederle Chorio- allantoic	Poxviridae	Vero	Selenazole (AVS-0253) and Arabinofuranosyladenine (AVS-1752)
Adenovirus Type 2	Adenoid 6	Adenoviridae	НЕр-2	Ribavirin (AVS-0001), Selenazole (AVS-0253)
Yellow Fever Virus	Asibi 17D (Prescreen)	Flaviviridae Flaviviridae	Vero Vero	Selenazole (AVS-0253) Selenazole (AVS-0253)
Japanese Encephalitis Virus	Nakayama	Flaviviridae	Vero	Selenazole (AVS-0253)
Venezuelan Equine Encephalomyelitis Virus	Trinidad Donkey	Togaviridae	Vero	Selenazole (AVS-0253)
Punta Toro Virus	Adames	Bunyaviridae	Vero	Ribavirin (AVS-0001)
Sandfly Fever Virus	Sicilian	Bunyaviridae	Vero	Ribavirin (AVS-0001)
Hantaan Virus	76-118	Bunyaviridae	Vero	Ribavirin (AVS-0001)
Pichinde Virus	4763	Arenaviridae	Vero	Ribavirin (AVS-0001)
Vesicular Stomatitis Virus	Indiana	Rhabdoviridae	1929	Carbocyclic-3-deaza-adenosine (AVS-0303)
Human Immunodeficiency Virus	IIIB	Retrovirus	ATH8,MT-2 CEM	3'-Azido-3'-deoxythymidine (AZT) (AVS-1603) 2',3'-Dideoxycytidine (ddC) (AVS-2639)

			E	XAMPLE	OF A	REPORT	OF MT	r assa	ΑY			
PLA	TE PW		11	N VITRO	ANTIVI	RAL RE	ESULTS				VS 0001	
DSDG	DEDG 0001		MTT ASSAY TAI: >30.3							>30.31	SI: >8.5	3
	1	2	3	4	5	6	7		9	10	11	12
A	0.126	0.122	0.138	0.122	0.114	0.116	0.039	0.039	0.039	0.039	0.040	0.040
	tox	00/10		0001 exports 0.305	0.276	1.514					00/10	
C	1.603	1.575	0.494	0.406	0.274	1.514					1.537	
n / 5	1.622	1.578	0.634	0.654	0.525	1.555					1.520	
\ =	1.578	0.502	1.017	0.999	1.021	1.616					0.523	
1 7	1.626	0.468	1.554	1.539	1.542	1.592					0.500	
	1.498	0.456	1.403	1.403	1.403	1.392					0.566	
			drug 0001 es		inground							
[2]	0.119	0.113	0.112	0.131	0.115	0.116	- highest dru					
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tion REAG	ENTE S CONTENCE CONTENCE		0.123 0.300 1.572 1.193	MAIAIM IC	: (mg/ml) : (mg/ml) ml imme: mal test		> 100, 5, > 18,	.48: .20:	DATE REAL 50% > 100 11 > 8	.00 .70 .53	05/01/89 95%	
view cent	ENTE S CONTENCE CONTENCE	DRUG ROW ON PLACE	0.123 0.380 1.572 1.193	APTIVII HEAD	(mg/ml) f (mg/ml) ml immer	ALUES VIRAL CPS	> 100; 5; > 18;	20 CYTOTOX HEAS	DATE REAL SOR > 100 LI > 8	.00: .70 .53 * VALUES * CELL	05/01/89 95% > 100 	GTRIC
view cent	ENTE S CONTENCE CONTENCE	DRUG ROW ON PLACE LOW B	0.123 0.380 1.572 1.193 0001 COMC. (uG/mL)	APTIVIS HEAP O.D. 137	(mg/ml) f (mg/ml) ml immer	VIRAL CPR 100%	> 100, 3: > 10.	AE: 20 CYTOTOX HEAF O.D. 1.443	DATE REAL SOR > 100 LI > 8	O	05/01/89 95% > 100 	GIRLC DITROL 007
view cent	ENTE S CONTENCE CONTENCE	DRUG ROW ON PLATE LOW B	0.123 0.380 1.572 1.193 0001 COMC. (uG/mL) 0.32	APPIVIA APPIVIA O.D. 137 117	(mg/ml) f (mg/ml) ml immer	VIRAL CPE 100%	> 100, 3: > 10.	20 CYTOTOX HEAF 0.D. 1.443 1.433	DATE REAL SOR > 100 LI > 8	CELL LABILITY 920	05/01/89 95% > 100 	GIRIC DITROL 007 008
view cent	ENTE S CONTENCE CONTENCE	DRUG ROW ON PLACE LOW B	0.123 0.380 1.572 1.193 0001 COMC. (uG/mL)	APTIVIS HEAP O.D. 137	(mg/ml) f (mg/ml) ml immer	VIRAL CPE 1009 1009	> 100, 3: > 18.	20 20 HEAF O.D. 1.443 1.433	DATE REAL SOR > 100 LI > 8	CELL LABILITY 92% 93%	05/01/89 95% > 100 	GIRLC DITROL 007
view cent	ENTE S CONTENCE CONTENCE	DRUG ROW ON PLACE LOW 2 C	0.123 0.380 1.572 1.193 0001 COMC. (ug/mL) 0.32 1	APPIVIA APPIVIA O.D. 137 117	(mg/ml) f (mg/ml) ml immer	VIRAL CPE 100%	> 100, 3: > 18.	20 CYTOTOX HEAR O.D. 1.443 1.433 1.458 1.458	DATE REAL SOR > 100 LI > 8	CELL LABILITY 920	05/01/89 95% > 100 	GETRIC DETROL 007 008 0.008
ion Div	ENTE S CONTENCE CONTENCE	DRUG ROW ON PLATE LOW 2 C	0.123 0.380 1.572 1.193 0001 COMC. (ug/mL) 0.32 1 3.2	APPIVIA APPIVI	(mg/ml) f (mg/ml) ml immer	7ALUES 1 VIRAL 1009 1009 929 569	> 100, 3. > 18.	20 20 HEAF O.D. 1.443 1.433	DATE REAL SOR > 100 LI > 8	CELL LABILITY 920 930	05/01/89 95% > 100 	007 008 0.008 011
view cent	ENTE S CONTENCE CONTENCE	DEUG ROW ON PLATE LOW B C D E	0.123 0.380 1.572 1.193 0001 COMC. (ug/mL) 0.32 1 3.2 10 32	AFFIVIN AFFIVIN MEAN 0.D. -117 0.094 0.521 1.053 0.905	: (mg/mg) : (mg/mg) hg Immus hg Immus h	7ALUES 7 VIRAL CPE 1009 1009 929 569 129 249	> 100, 3. > 18.	AB 200 CYTOTOX MEAN O.D. 1.443 1.458 1.485 1.496 1.326	SON	0 00% 270 233 2 VALUES 3 CELL 1ABILITY 923 913 944 955	05/01/89 95% > 100 	GTRIC DETROL 007 008 0.008 011 010

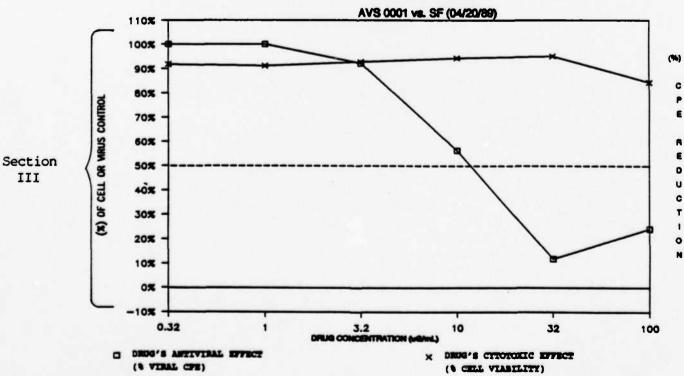


Figure 1

EXPLANATION OF IN VITRO ANTIVIRAL RESULTS FORM (MTT ASSAY)

The In Vitro Antiviral Results form (Figure 1) has three sections:

Section I. Sample and Test Identification and the actual raw data (optical density readings) collected for each 96-well plate.

The Test Identification and Raw Data Section specifies the compound that was tested, a unique plate number (assigned by the computer) and the actual optical density readings for the virus control (vc), cell control (cc), drug alone (tox) and the drug plus virus (drug experimental). Background readings are also taken for the container (plastic), reagent (culture medium) and drug colorimetric.

Section II. Printouts of pertinent Control and Test Results computed from the raw data are in this section.

Tabular Dose Response Test Results are calculated as follows:

- a. Mean Medium O.D. is subtracted from means of virus and cell control O.D.s.
- b. Drug colorimetric O.D. is subtracted from each infected and uninfected value at the corresponding drug dilution.
- c. The Differential is the mean O.D. attributable to virus kill (CPE).

Differential = (mean O.D. cell control) - (mean O.D. Virus control).

d. For infected cells at each drug concentration,

% CPE = the reciprocal of:

[(mean O.D. infected wells - mean O.D. drug colorimetric wells) -(mean O.D. virus control - mean reagent O.D.)]/Differential; expressed in percent.

e. For uninfected cells at each drug dilution,

% Cell Viability = [(mean O.D. uninfected drug-treated wells) - (mean drug colorimetric wells)]/(mean O.D. cell control).

Quantitation of Viral Cytopathic Effect (CPE) and Drug Activity are displayed in the shaded area of Section II. These values are defined below:

 $TC_{25,50,95}$ = The drug concentration ($\mu g/ml$) that reduced cell viability by 25%, 50% or 95%.

 $IC_{25,50,95}$ = The drug concentration (μ g/ml) that inhibited CPE by 25%, 50% or 95% calculated by using a regression analysis program for semilog curve fitting.

 $AI_{25,50,95}$ = Antiviral Index, calculated by dividing $TC_{25,50}$ or $_{95}$ by $IC_{25,50}$ or $_{95}$.

TAI = Total Antiviral Index - the area between the cytotoxicity and the antiviral curves.

SI = Selectivity Index, calculated by dividing the TC₂₅ by the IC₅₀.

Figure 1 (Cont'd)

Section III. The Graphic Results Summary Section displays a plot or graphic illustration from computed values in Section II.

The line connecting the square symbols depicts the percentage of viral CPE in virus-infected cells treated with the test compound (at the indicated concentrations) relative to the Differential. This line expresses the *in vitro* anti-viral activity of the sample.

The line connecting the X symbols depicts the percentage of surviving uninfected cells treated with the test compound relative to the uninfected, untreated control (cell control). This line expresses the cytotoxicity of the drug at the various concentrations, or percent cell viability. The dotted line is just a reference line at 50%.

3.3.1 Vaccinia Virus (VV)

Vaccinia virus, strain Lederle CA, was obtained from Dr. Wilton Rightsel, formerly with Parke, Davis and Company, Detroit, Michigan. We have serially passed VV in HEp-2 (continuous-passage human carcinoma of the larynx) and Vero (continuous-passage African green monkey kidney) cell monolayer cultures. The VV used to screen compounds for USAMRIID was propagated and assayed in Vero cell monolayer cultures in Eagle's Minimal Essential Medium (MEM) supplemented with 2% hear-inactivated fetal bovine serum (Δ fbs) and 50 μ g/ml of gentamicin. Virus stocks were titrated according to the procedure of Reed and Muench (1938) and diluted in culture medium to 100 CCID₅₀ per 0.1 ml.

During the time period covered by this report, compounds were screened for activity against VV in Vero cell monolayer cultures in 96-well plates, employing a CPE-inhibition assay procedure.

Subsequent to 12/15/88 and beginning with Shipment 42, all compounds have been screened for activity against VV by the MTT assay procedure, in which the degree of cell viability (and therefore CPE and drug cytotoxicity) is determined by MTT uptake. This procedure is based on the reduction of the tetrazolium salt, 3-(4,5-dimethyl-thiazol-2-yl)-2,5-diphenyl tetrazolium bromide (MTT) by mitochondrial enzymes of viable host cells to MTT formazan. The blue color of the MTT formazan is measured spectrophotometrically. Vero cells are seeded into 96-well plates at a density of 4 x 10⁴ cells in 0.2 ml per well in MEM + 5% bovine calf serum. The plates are incubated at 37°C overnight. The next day our CPE-inhibition assay is set up according to the format shown in Appendix A. MEM + 2% heatinactivated fetal bovine serum (\Delta fbs) serves as experiment medium. To each of triplicate test wells containing replicate cell monolayers, 100 µl of each test drug solution (or suspension) and 100 µl of experiment virus are dispensed. Six 0.5 \log_{10} concentrations of each drug beginning with 320 μ g/ml are routinely used. Compounds are solubilized or suspended and diluted in culture medium the day of use. Cell controls containing 200 µl of medium, virus controls containing cells, medium and virus, and duplicate drug cytotoxicity controls containing cells, medium and each drug concentration are included on each plate. Blank wells, medium (Reagent) control wells, and drug colorimetric controls (drug + medium + MTT + SDS - no cells) accompany each test. The covered plates are incubated at 37°C in a humidified atmosphere containing 2% CO₂.

When CPE reach 100%, 5 days postvirus infection (p.i.), 20 μ l of MTT (a 5 mg/ml solution in PBS) are added to each well. The plates are incubated at 37°C for six to seven hours to allow reduction of the MTT to the formazan form. Then 40 μ l aliquots of a 30% solution of SDS in 0.02N HCl are added to the plate wells. The plates are incubated overnight to allow the SDS to lyse the cells and dissolve the MTT formazan crystals. The absorbance of the contents of each well is determined by a plate reader employing dual filters of 570 and 650 nm. The plate reader is interfaced with a computer programmed to capture the optical density (O.D.) measurements from the reader and calculate indices such as the IC_{25, 50, 95}, TC_{25, 50, 95} and to plot the percents of viral CPE and cell viability of drug-treated cultures.

3.3.2 Adenovirus Type 2 (AD2)

We obtained the Adenoid-6 strain of Adenovirus Type 2 from the American Type Culture Collection. In our laboratories, the virus was passaged in HEp-2 cell monolayer cultures in MEM supplemented with 0.4% lactalbumin hydrolysate and 2% heat-inactivated fbs. An early "cytotoxicity" is frequently observed in adenovirus-infected cell cultures. This "cytotoxicity", the result of penton toxin following an accumulation of unassembled viral components within the cell, can mask the later-appearing CPE. To prevent the appearance of this early cytotoxicity, virus stocks were treated as follows: Adenovirus-infected HEp-2 cell cultures were subjected to three cycles of freeze-thawing to disrupt the cell membranes. The material from the cultures was centrifuged, and the pooled supernatant fluid was treated with 100 µg/ml of purified trypsin (Worthington, TRL) for one hour at 37 °C. The enzyme

action was then stopped by adding soybean trypsin inhibitor (Worthington; titrated for specific activity against trypsin) to the supernatant viral fluid. The trypsin-treated virus preparation, along with the untreated supernatant viral fluid was titrated for infectivity in HEp-2 cell cultures to make sure that the viral infectivity of the treated preparation was not diminished. Before the trypsin-treated stock adenovirus was used for antiviral screening, it was evaluated as the challenge virus against our positive control compounds Selenazole and Ribavirin.

During the period covered by this report, test compounds were screened for antiviral activity against adenovirus using a CPE-inhibition assay procedure as described above for vaccinia virus in Section 3.3.1. Stock AD2 was diluted in medium to 100 CCID₅₀ per 0.1 ml. We have found that a 32-CCID₅₀ challenge of AD2 per plate well resulted in only 50-70% CPE in four days. Because HEp-2 cells grow rapidly, the host cell cultures in the plate wells deteriorated significantly after four days post-virus infection. Therefore the stronger virus challenge allowed 100% CPE to develop in four days. Six 0.5 log₁₀ concentrations of each compound were evaluated in triplicate HEp-2 cell culture wells. To each culture well was added 0.1 ml of the test drug solution, followed immediately by 0.1 ml of the challenge virus dilution. The 96-well plates were incubated for four days at 37°C in a humidified atmosphere containing 2% CO₂. The CPE were examined microscopically and graded 0-4.

As we discussed in our Sixth Semiannual Progress Report (SRI-89-BIO-109-5975, December 15, 1988) an MTT CPE-inhibition assay was developed for use with AD2 using A549 cells or 293 cells. The MTT assay in either cell line was dose-responsive and reproducible, but would be a more stringent screening assay than the present CPE-inhibition assay which is read microscopically. Therefore we continued using the CPE-inhibition (microscopic) assay in HEp-2.

As a result of contract modifications by USAMRIID in November, 1989, AD2 was deleted from the virus spectrum utilized for primary screening.

3.3.3 Yellow Fever Virus (YF)

We obtained our original stock of Yellow Fever Virus, Asibi strain, from Dr. Andrew J. Main, Jr., of the Yale Arbovirus Research Unit, New Haven, Connecticut.

To grow virus stocks, Vero cells (ATCC) were infected at an moi of about 0.1 PFU/cell in MEM containing 10% inactivated fbs. Virus was allowed to adsorb for 1 hour, after which a minimal volume of growth medium was added and the cells were incubated at 37°C. The culture fluid was collected at 5 days post infection and the clarified by centrifugation (5000 rpm, 15 minutes, 4°C) in a Sorvall SA-600 rotor. The supernate was dispensed into 0.5 ml aliquots and then frozen and stored at -84°C. One aliquot was used to determine the TCID₅₀ and PFU titers for the stock.

The standard operating procedure is an MTT-based assay as follows:

Vero cells are seeded in 96-well tissue culture plates at a density of 1.8 x 10^4 cells/well in 100 μ l complete EMEM and incubated overnight at 37°C/5% CO₂.

The following day, the medium is aspirated out of the wells and 50 μ l of virus (diluted to a virus load of 32 TCID₅₀/well) are added to the test and virus control wells. The toxicity and cell control wells receive 50 μ l of complete EMEM. The plates are incubated for 30 minutes at 37°C/5% CO₂ to allow virus to adsorb.

After virus adsorption, the test wells receive an additional 50 μ l of a 2X concentration of drug and the control wells receive an additional 50 μ l of complete EMEM. The total volume per well = 100 μ l. Generally, the standard drug test concentration range is from 1 μ g/ml to 320 μ g/ml.

The plates are incubated at 37°C/5% CO₂ until the virus control wells exhibit maximum CPE (+4 = 100%) by visual scoring under the microscope (40X). Generally, the YF virus (with 32 TCID₅₀ virus load) requires 6-7 days of incubation to achieve maximum CPE. At this time, all wells (excluding plastic control wells) receive 50 μ l of MTT solution (1.50) mg/ml, prepared in complete EMEM).

The plates are incubated 7 hours at 37°C/5\% CO₂ (in the dark) to allow reduction of the tetrazolium salt to the formazan product, after which the cells are lysed and the formazan is solubilized by the addition of $100 \,\mu\text{l}$ of a lysing solution (10% SDS-0.01N HCl) to all wells generally. If the MTT solution is added in the morning, the lysing agent is added at the end of the day, and lysis occurs overnight (14 - 15 hours in the dark).

3.3.4 Japanese Encephalitis Virus (JE)

We obtained our original stock of Japanese Encephalitis Virus, Nakayama strain, from Dr. George R. French of the Salk Institute, Swiftwater, Pennsylvania.

To grow virus stocks, Vero cells (ATCC) were infected at an moi of about 0.1 PFU/cell in MEM containing 5% fbs. Virus was allowed to adsorb for 1 hour, then a minimal volume of growth medium was added and the cells were incubated at 37°C. The culture fluid was collected at 4-5 days post infection and clarified by centrifugation (5000 rpm, 15 minutes, 4°C) in a Sorvall SA-600 rotor. The supernate was dispensed into 0.5 ml aliquots and then frozen and stored at -84°C. One aliquot was used to determine TCID₅₀ and PFU titers of the stock.

The standard operating procedure is an MTT-based assay as follows:

Vero cells are seeded in 96-well tissue culture plates at a density of 1.8 x 10^4 cells/well in 100 μ l complete EMEM and incubated overnight at 37°C/5% CO₂.

The following day, the medium is aspirated out of the wells and 50 μ l of virus (diluted to a virus load of 32 TCID₅₀/well) are added to the test and virus control wells. The toxicity and cell control wells receive 50 μ l of complete EMEM. The plates are incubated for 30 minutes at 37°C/5% CO₂ to allow virus to adsorb.

After virus adsorption, the test wells receive an additional 50 μ l of a 2X concentration of drug and the control wells receive an additional 50 μ l of complete EMEM. The total volume per well = 100 μ l. Generally the standard drug test concentration range is from 1.0 μ g/ml to 520 μ g/ml.

The plates are incubated at $37^{\circ}\text{C}/5\%$ CO₂ until the virus control wells exhibit maximum CPE (+4 = 100%) by visual scoring under the microscope (40X). Generally, the JE virus (with 32 TCID₅₀ virus load) requires 6-7 days of incubation to achieve maximum CPE. At this time, all wells (excluding plastic control wells) receive 50 μ l of MTT solution (1.50) mg/ml, prepared in complete EMEM).

The plates are incubated 7 hours at $37^{\circ}\text{C}/5\%$ CO₂ (in the dark) to allow reduction of the tetrazolium salt to the formazan product, after which the cells are lysed and the formazan is solubilized by the addition of 100 μ l of a lysing solution (10% SDS-0.01N HCl) to all wells. If the MTT solution is added in the morning, the lysing agent is added at the end of the day, and lysis occurs overnight (14 - 15 hours in the dark).

3.3.5 <u>Venezuelan Equine Encephalomyelitis Virus (VE)</u>

We obtained our original stock of Venezuelan Equine Encephalomyelitis Virus, Trinidad Donkey strain, from Dr. George R. French of The Salk Institute, Swiftwater, Pennsylvania.

The virus was propagated in Vero cells (ATCC) by infection at a moi of < 0.1 PFU/cell. Fluids from infected cultures were collected three days post-infection and clarified by centrifugation (5000 rpm, 15 min, 4°C) in a Sorvall SA-600 rotor. The supernatant fluid was dispensed into 0.5-ml aliquots, quick frozen in a dry ice-ethanol bath and stored at -84°C. One aliquot was used to determine TCID₅₀ and PFU titers of the stock.

The standard operating procedure is an MTT-based assay as follows:

Vero cells are seeded in 96-well tissue culture plates at a density of 1.8 x 10^4 cells/well in 100 μ l complete EMEM and incubated overnight at 37°C/5% CO₂.

The following day, the medium is aspirated out of the wells and 50 μ l of virus (diluted to a virus load of 32 TCID₅₀/well) are added to the test and virus control wells. The toxicity and cell control wells receive 50 μ l of complete EMEM. The plates are incubated for 30 minutes at 37°C/5% CO₂ to allow virus to adsorb.

After virus adsorption, the test wells receive an additional 50 μ l of a 2X concentration of drug and the control wells receive an additional 50 μ l of complete EMEM. The total volume per well = 100 μ l. Generally the standard drug test concentration range is from 1.0 μ g/ml to 320 μ g/ml.

The plates are incubated at $37^{\circ}\text{C}/5\%$ CO₂ until the virus control wells exhibit maximum CPE (+4 = 100%) by visual scoring under the microscope (40X). Generally, the VEE (with 32 TCID₅₀ virus load) requires 3 days of incubation to achieve maximum CPE. At this time, all wells (excluding plastic control wells) receive 50 μ l of MTT solution (1.50) mg/ml, prepared in complete EMEM).

The plates are incubated 7 hours at $37^{\circ}\text{C}/5\%$ CO₂ (in the dark) to allow reduction of the tetrazolium salt to the formazan product, after which the cells are lysed and the formazan is solubilized by the addition of 100 μ l of a lysing solution (10% SDS-0.01N HCl) to all wells. If the MTT solution is added in the morning, the lysing agent is added at the end of the day, and lysis occurs overnight (14-15 hours in the dark).

3.3.6 Punta Toro Virus (PT)

We obtained our original stock of Punta Toro virus (Adames strain) from Dr. Robert Sidwell of Utah State University. His virus stock originated from Dr. Dominique Pifat of USAMRIID.

To grow virus stocks, Vero cells (ATCC) were infected at a low multiplicity of infection (moi; ≤ 0.1) in MEM supplemented with 10% inactivated fbs. Culture fluid was collected at five days post-infection and clarified by centrifugation (5000 rpm, 15 minutes, 4°C) in a Sorvall SA-600 rotor. The supernate was dispensed into 0.5-ml aliquots, then frozen and stored at -84°C. One aliquot was used to determine a TCID₅₀ titer and a plaque forming unit (PFU) titer for the stock.

The standard operating procedure is an MTT-based assay as follows:

Vero cells are seeded in 96-well tissue culture plates at a density of 1.8 x 10^4 cells/well in 100 μ l complete EMEM and incubated overnight at 37°C/5% CO₂.

The following day, the medium is aspirated and all cells are treated with 100 μ l of pretreatment solution for 30 minutes at 37°C/5% CO₂. The pretreatment solution is composed of 1% DMSO from a 100% DMSO stock solution, 1% DEAE-Dextran from a 2.0 mg/ml stock solution in sterile water, and 98% Hank's Balanced Salt Solution.

After this the pretreatment medium is aspirated out of the wells and 50 μ l of virus (diluted to a virus load of 32 TCID₅₀/well) are added to the test and virus control wells. The toxicity and cell control wells receive 50 μ l of complete EMEM. The plates are incubated for 30 minutes at 37°C/5% CO₂ to allow virus to adsorb.

After virus adsorption, the test wells receive an additional 50 μ l of a 2X concentration of drug and the control wells receive an additional 50 μ l of complete EMEM. The total volume per well = 100 μ l. Generally the standard drug for concentration range is from 1 μ g/ml to 320 μ g/ml.

The plates are incubated at 37°C/5% CO₂ until the virus control wells exhibit maximum CPE (+4 = 100%) by visual scoring under the microscope (40X). Generally, the PT virus (with 32 TCID₅₀ virus load) requires 6-7 days of incubation to achieve maximum CPE. At this time, all wells (excluding plastic control wells) receive 50 μ l of MTT solution (1.50) mg/ml, prepared in complete EMEM).

The plates are incubated 7 hours at $37^{\circ}\text{C}/5\%$ CO₂ (in the dark) to allow reduction of the tetrazolium salt to the formazan product, after which the cells are lysed and the formazan is solubilized by the addition of 100 μ l of a lysing solution (10% SDS-0.01N HCl) to all wells. Generally if the MTT solution is added in the morning, the lysing agent is added at the end of the day, and lysis occurs overnight (14 - 15 hours in the dark).

3.3.7 Sandfly Fever Virus (SF)

We obtained our original stock of Sandfly Fever virus, Sicilian strain (TC adapted), from Dr. George R. French of the Salk Institute, Government Services Division, Swiftwater, Pennsylvania.

To grow virus stocks, Vero-76 cells (Dr. French, Salk Institute) or Vero cells (ATCC), depending upon availability, were pre-treated with DEAE-dextran (25 μ g/ml) and 1% DMSO in growth medium (MEM containing 10% inactivated fbs) for 30 minutes at 37°C. This was removed and then the cells were infected with SF virus in growth medium at an moi of 0.1. Virus was allowed to adsorb for one hour at 37°C. A minimal volume of growth medium was then added and the cells were incubated at 37°C. The culture fluid was collected at four days post infection and clarified by centrifugation (5000 rpm, 15 min, 4°C). The supernate was dispensed into 0.5 ml aliquots, then frozen and stored at -84°C. One aliquot was used to determine TCID₅₀ and PFU titers for the stock.

The standard operating procedure is an MTT-based assay as follows:

Vero cells are seeded in 96-well tissue culture plates at a density of 1.8 x 10^4 cells/well in 100 μ l complete EMEM and incubated overnight at 37°C/5% CO₂.

The following day, the medium is aspirated and all cells are treated with $100 \mu l$ of pretreatment solution for 30 minutes at $37^{\circ}\text{C}/5\%$ CO₂. The pretreatment solution is composed of 1% DMSO from a 100% stock solution, 1% DEAE-Dextran from a 2.5 mg/ml stock solution in sterile water, and 98% HBSS.

After this, the pretreatment medium is aspirated out of the wells and 50 μ l of virus (diluted to a virus load of 32 TCID₅₀/well) are added to the test and virus control wells. The toxicity and cell control wells receive 50 μ l of complete EMEM. The plates are incubated for 30 minutes at 37°C/5% CO₂ to allow virus to adsorb.

After virus adsorption, the test wells receive an additional 50 μ l of a 2X concentration of drug and the control wells receive an additional 50 μ l of complete EMEM. The total volume per well = 100 μ l. Generally the standard drug concentration range is from 1 μ g/ml to 320 μ g/ml.

The plates are incubated at $37^{\circ}\text{C}/5\%$ CO₂ until the virus control wells exhibit maximum CPE (+4 = 100%) by visual scoring under the microscope (40X). Generally, the SF virus (with 32 TCID₅₀ virus load) requires 6-7 days of incubation to achieve maximum CPE. At this time, all wells (excluding plastic control wells) receive 50 μ l of MTT solution (1.50) mg/ml, prepared in complete EMEM).

The plates are incubated 7 hours at $37^{\circ}\text{C}/5\%$ CO₂ (in the dark) to allow reduction of the tetrazolium salt to the formazan product, after which the cells are lysed and the formazan is solubilized by the addition of 100 μ l of a lysing solution (10% SDS-0.01N HCl) to all wells. Generally if the MTT solution is added in the morning, the lysing agent is added at the end of the day, and lysis occurs overnight (14 - 15 hours in the dark).

3.3.8 Hantaan Virus (HTN)

We obtained several vials of Hantaan Virus, strain 76-118, (formerly referred to as Korean Hemorrhagic Fever Virus, KHF) from Dr. John W. Huggins and Mr. Orville Brand of USAMRIID. Initial attempts to plaque this virus in a routine plaque assay were unsuccessful. Discussions with Dr. Huggins and Mr. Brand indicated that the quality of the reagents used in the assay is critical. Mr. Brand graciously supplied us with a shipment of his tested cell culture supplies and fresh cultures of Vero VSC-6 cells to use in assaying KHF virus.

Six-well culture trays of VSC-6 cells were seeded at a density of 3.75×10^5 cells/well with MEM containing 5% horse serum, 5% Nu serum, 10 mM HEPES buffer, 2 mM glutamine, 1 mM sodium pyruvate, and penicillin/streptomycin (at 100 units and 100 μ g/ml, respectively) and incubated at 37°C in a 5% CO₂ humidified atmosphere. As suggested by Mr. Brand, KHF virus was diluted ten-fold in MEM with supplements and stored at 4°C and stored overnight prior to assay.

The following day, the culture medium was aspirated and 0.4 ml of virus dilution was added to duplicate cultures. Two cultures per tray were mock-infected with 0.4 ml MEM. Culture trays were incubated for 1 hour at 37°C and swirled every 15 minutes.

After 1 hour adsorption, 2.5 ml of overlay consisting of cell culture medium plus 0.6% agarose was added to each culture. The trays were reincubated at 37°C. Seven days after infection, a 2.0 ml volume of overlay was added to each culture and the cultures reincubated. Twelve days after infection, 1 ml of overlay with neutral red was added to each well. Culture trays were reincubated and observed for plaques from Day 13 to Day 20, post infection.

Testing of antiviral compounds against the Hantaan virus was temporarily suspended in our laboratory to complete the development of the ELISA system then testing was carried out at USAMRIID Ft. Detrick, Maryland under the direction of Dr. John Huggins.

3.3.9 Pichinde Virus (PIC)

Vero cells, obtained from Dr. George R. French of The Salk Institute, Swiftwater, Pennsylvania, were maintained in Eagle's Minimal Essential Medium containing 10% fetal bovine serum. Baby hamster kidney (BHK-21) cells were obtained from the American Type Culture Collection and maintained in MEM containing 10% newborn calf serum and 10% tryptose phosphate broth.

Pichinde virus strain 4763 was obtained from Dr. J. Gangemi of the University of South Carolina, School of Medicine. The virus preparation received was that of a guinea pig spleen homogenate from Pichinde virus infected guinea pigs. After amplifying this virus preparation in BHK-21 cells, a three-time plaque-purified virus preparation was generated in Vero cells. The plaque-purified virus was amplified twice in BHK-21 cell cultures and stored at -84°C as a virus seed stock. Several

preparations of virus used for plaque-reduction assays were prepared by infecting BHK-21 cells with seed stock virus at a multiplicity of infection of 1 PFU/cell. Culture fluids collected three days post-infection were clarified by centrifugation at 600 x g for 5 min., frozen rapidly in a dry ice/ethanol bath and stored at -84°C in small aliquots. These preparations had virus titers ranging from 6.4 x 10⁷ to 1.2 x 10⁹ PFU/ml.

Vero cells, seeded into six-well cell culture trays at a density of 8 x 10^5 cells per well, were incubated at 37°C in a humidified 5% CO_2 incubator. The following day the cultures were examined microscopically for confluency. Ribavirin and experimental antiviral drugs were serially diluted in half- log_{10} increments in MEM containing 10% fbs. The final test concentration range was from 1 - $320\mu g/ml$. Culture fluids were removed by aspiration and to four of the six cultures of each tray were added a 0.2 ml inocula of PIC virus diluted in MEM containing 10% fbs so as to contain 40 to 80 PFU. To the remaining cultures of each tray, 0.2 ml of culture medium alone was added. The cultures were reincubated as above and periodically shaken so that the inoculum was dispensed over the entire monolayer.

One hour later, 0.2 ml of the appropriate dilution of drug was added to three of the virus-infected cultures (virus-infected, treated) and to one of the mock-infected cultures (toxicity control) on each tray. To the remaining mock-infected (cell control) and virus-infected (virus control) cultures were added 0.2 ml of MEM containing 10% fbs. The culture fluids were mixed by shaking the culture trays, then 1.6 ml of agar overlay was added and each culture was thoroughly mixed as above. The cultures were set on a level surface until the agar overlay solidified and were then reincubated at 37°C. The agar overlay used in the assay contained Eagle's MEM, fetal bovine serum, glutamine, non-essential amino acids, sodium bicarbonate, and Noble agar in final concentrations of 1x, 2%, 1 mM, 1mM, 2.2 mg/ml and 1%, respectively.

Three days later 2 ml of a sterily filtered neutral red solution (0.08 mg/ml in PBS) was added to each culture. Cultures were reincubated; then six to eight hours later plaques were examined and counted. With each daily assay, Ribavirin (positive control antiviral drug) was included in a parallel assay. Drug cytotoxicity was microscopically evaluated and rated as with CPE-inhibition assays.

3.3.10 Vesicular Stomatitis Virus (VSV)

The VSV (Indiana strain) was obtained from Dr. Robert Sidwell, Utah State University, Logan, Utah in 1973. Dr. Sidwell had originally obtained the virus from the ATCC. Since the virus has been in this laboratory, it has been passaged 5 times in L929 cells.

The L929 cells were obtained from the ATCC and have been carried in continuous passage or stored frozen in liquid nitrogen. The medium used to maintain the L929 cells was Eagle's minimum essential medium supplemented with 10% or 5% heat-inactivated fetal bovine serum, 100 units penicillin per ml, and $100 \mu g$ streptomycin per ml (EMEM + 10% or 5% heat-inactivated fbs).

From November 16, 1988 to January 13, 1989, CPE-inhibition assay was used to assay the drugs against VSV. The L929 cells were seeded into the 96-well plates at a concentration of 4×10^4 cells per well in 0.2 ml medium. The cells were pregrown for 20 hrs. before the addition of drug and virus.

Stock virus was pretitered according to the method of Reed and Muench (1938) and diluted in cell culture medium to yield 32 $TCID_{50}$ (tissue culture infectious dose, 50%) units per 0.1 ml. Antiviral assays were designed to test seven 0.5- log_{10} concentrations of each compound in triplicate against the challenge viruses. If drug cytotoxicity was provided, then drug concentrations were chosen to range from toxic to noncytotoxic. If no cytotoxicity information was available, testing began at 320 μ g/ml. To each

of the replicate host cell cultures was added 0.1 ml of the test drug solution and 0.1 ml of virus suspension. Cell controls containing medium alone, virus controls containing medium and virus, and drug cytotoxicity controls containing medium and each drug concentration were included on each plate. The plates were incubated at 37°C and virus-induced CPE were scored on Day 3 post-virus infection.

The degree of inhibition of virus-induced CPE and the degree of drug-induced cytotoxicity were observed microscopically. CPE were scored numerically from 0 (normal uninfected cells) to 4 (100% virus-induced cell destruction) for each individual culture as follows:

- 4 = Viral CPE observed in 100% of the cells
- 3 = Viral CPE observed in 75% of the cells
- 2 = Viral CPE observed in 50% of the cells
- 1 = Viral CPE observed in 25% of the cells
- 0 = No CPE; normal cell monolayer
- u = Unsatisfactory test, e.g., contamination or leakage
- t = Drug is toxic to cells, CPE not discernible
- p = Drug is partially toxic to cells, cell monolayer is intact so that CPE may be discernible

Antiviral activity was determined by calculating the degree of inhibition of virus-induced CPE in drug-treated, virus-infected cell cultures by means of a virus rating (VR). The VR is a standard weighted measurement of antiviral activity taking into account both the degree of CPE inhibition and drug cytotoxicity, and was determined by a modification of the method of Ehrlich et al. (1965). The VR was calculated as 0.1 of the sum of the numerical differences between the recorded CPE grade of each test well and that of the corresponding virus control culture. Numerical differences between the reading of test wells containing a drug concentration which was partially cytotoxic (p) and their corresponding virus controls were halved. As requested by USAMRIID, the VR was also reported by the method of Sidwell and Huffman (1971). This virus rating (VR*) was calculated by dividing the VR obtained by the method of Ehrlich, et al. (1965) by 3. The inhibitory drug concentration which reduced the CPE by 50% (ID₅₀) was calculated by using a regression analysis program for semilog curve fitting. A therapeutic index (TI) for each active test compound for each susceptible virus was determined by dividing the minimum cytotoxic concentration (MTC) of the test compound by the ID₅₀.

From January 14, 1989 to November 15, 1989, the following procedure was used to evaluate drugs in the MTT assay. L929 cells were seeded into 96-well tissue culture plates at a concentration of 4 x 10^4 cells/well on the day before the cells were used for the assay. A volume of 50 μ l of the appropriate drug concentration and 50 μ l of a virus suspension containing 32 CCID₅₀ of virus were added per well. On Day 3 after virus infection when CPE was complete in the virus controls, 50 μ l of a solution containing 2 mg/ml MTT in Eagle's MEM + 5% heat-inactivated fetal bovine serum was added. The cultures were incubated with MTT for 7 hours before addition of 100 μ l of 10% SDS:0.01N HCl per well. The cultures were incubated overnight and the following day the optical density was read at 570 nm on a Perkin-Elmer Lambda plate reader.

3.3.11 In Vitro Antiviral Screening: Retroviruses:

3.3.11.1 Primary Screening-HIV

3.3.11.1.1 Modified Broder assay

We obtained the human OKT4⁺ T-cell clone, ATH8, from Dr. Samuel Broder's laboratory (NCI) through Dr. Hiroaki Mitsuya. This cell line is quite sensitive to the cytopathic effect of Human Immunodeficiency Virus (HIV). We propagated these cells in RPMI 1640 medium supplemented with

4 mM glutamine, 15% heat-inactivated fetal bovine serum, antibiotics (50 units of penicillin and 50 μ g of streptomycin per ml), and 50 units of recombinant-derived human interleukin-2 (ala-125; AMGen Biologicals) per ml. The clone H9 cell line, an OKT4⁺ human T-cell line that is permissive for HIV replication but largely resistant to virus-induced CPE, H9 cells productively infected with the HTLV-IIIB strain of HIV, and H9 cells productively infected with the RF-II Haitian variant of HIV were obtained from Dr. Robert Gallo's laboratory (NCI) through Dr. Howard Streicher. We propagated these cell lines in RPMI 1640 medium supplemented with 4mM glutamine, 20% heat-inactivated fetal bovine serum, and antibiotics (50 units of penicillin and 50μ g of streptomycin per ml). For infectious virus we used undiluted culture supernates form H9/HTLV-IIIB producer cells. We have found that supernate collected 48 hours post-cell passage shows the best infectivity.

The initial screening of all compounds received from USAMRIID for retrovirus antiviral activity was done using the modified CPE-inhibition assay developed by Broder and co-workers (Mitsuya et al., 1985 and 1986). This assay is based on the ability of uninfected ATH8 cells to grow and form a pellet at the bottom of a culture tube. Starting about 4 days after HIV addition, infected ATH8 cells began to die and the pellet started to break up. The cell pellet was completely destroyed within 10 days. The protective effect of test compounds was assessed by adding them at varying concentrations to the cultured cells at the time of virus infection, then monitoring the status of the cell pellet.

ATH8 cells were used as the primary target cells in the HIV-induced CPE-inhibition assay. Cells were treated with polybrene ($2\mu g/ml$ in growth medium) for 30 minutes at 37°C, then collected by gentle centrifugation (40 xg for 15 minutes at room temperature) and resuspended in clarified (8000 xg for 15 minutes at 4°C) supernate freshly harvested form 48 hour post-passage H9/HTLV-IIIB cells. Following a 60-minute adsorption period at 37°C, the cells were dispensed into the U-bottom wells of 96-well trays (2 x 10^4 cells in 0.1 ml per well). An equal volume (0.1 ml) of supplemented RPMI 1640 medium containing test compound and twice the normal concentration of interleukin-2 was then added to each well. Test compounds were evaluated at seven half-log₁₀ dilutions. Triplicate virus-infected cultures and one uninfected compound cytotoxicity control culture were included at each dosage level. Cultures were incubated at 37°C in a humidified atmosphere of 5% CO₂ in air. The sizes of the cell pellets in the test compound wells were compared to the pellet sizes of infected and uninfected cell control wells daily for 10 days. On day 10 post-infection aliquots were taken from selected individual wells and the total cell number and cell viability (based on trypan-blue dye-exclusion) were determined. Dideoxycytidine was included as a positive control with each set of compounds tested.

3.3.11.1.2 XTT Assay

In order to accommodate the large volume of screening required by the NCI modification of this contract, an automated assay system was adapted based on the screening program in use by the NCI Developmental Therapeutics Program. We have used the human T-cell line MT-2 (Harada et al., 1985) in these assays. These cells have growth characteristics resembling transformed cells, do not require interleukin, and undergo a lytic infection with HIV. They are therefore well suited to a large scale screening program.

This assay resembles the modified Broder assay in the virus and drug inoculation regimens. At the time of assay analysis, however, the XTT assay involves addition of a tetrazolium dye (XTT) which is converted by viable cells to a soluble formazan derivative which is measured by optical density (O.D.). The O.D. value obtained for a given well is proportional to the number of viable cells in the well and is therefore a measure of virus-induced CPE (and also cytotoxicity) or conversely, the inhibition of CPE by a test compound.

As developed by the NCI (Developmental Therapeutics Program), the optical densities were read by an automated plate reader which is interfaced to a computer. In addition to collecting and analyzing

the test data, the computer system maintained schedules and plate identification numbers. The computer in the testing laboratory was linked to a unit in the compound preparation laboratory and both were on line to a mainframe computer at the NCI.

Assays were scheduled for testing using the ASGN module of the IVSS. We have begun specific drug assignments rather than utilizing the priority designation system from the mainframe Master files as we did at the beginning of the program. We changed this method primarily because the AVAIL module of the IVSS system, as written, cannot differentiate varying cell line assignments and the *in vitro* screening program now uses 2 cell lines, MT-2 and CEM.

In addition to AVAIL and ASGN which are accessed under MS-DOS on the IBM-370, there are XENIX operating system programs stored on a micro-computer in the treatment laboratory. These XENIX modules of the IVSS include Plate inoculation, Drug Addition, Stain Addition, and Plate Reading. The plate reading module uses a V MAX Kinetic Micro plate reader interfaced to the XENIX program on the micro-computer. Data from this computer is automatically up-loaded and down-loaded from the IVSS system managed by VSE in Alexandria, VA. Plate Analysis Reports are automatically generated and printed locally during Plate Read before automatic up-load of the data to the mainframe.

Requests for additional quantities of drugs to be retested, and periodic over-all status reports were communicated to DTP staff primarily through electronic mail services on the NIH mainframe computers.

3.3.11.2 Secondary Evaluations

3.3.11.2.1 Feline Leukemia Virus - FAIDS Variant (FeLV)

We obtained our initial stock of Feline Leukemia Virus, FAIDS variant from Dr. Ed Hoover of Colorado State University.

Following the procedure for growing stock virus established by Dr. Hoover, feline embryonic fibroblasts, AH-927 (Dr. Hoover), were grown in MEM containing 10% inactivated FBS and 1% non-essential amino acids. Twenty-four hours prior to infecting with virus, the cells were subcultured at a ratio of 1:4 and 0.2ml of Polybrene (2 mg/ml stock) was added for each 100 ml growth medium. On the day of infection, the growth medium was removed and the cells were washed twice with PBS. The virus inoculum was allowed to adsorb for 1 hour at 37°C. The virus inoculum was removed, the cells washed twice with PBS, and then a minimal volume of growth medium containing Polybrene was added. After five days incubation the culture fluid was collected and clarified by centrifugation (5000 rpm x 15 min). The supernate was dispensed into 1.0-ml aliquots, frozen, and then stored at -120°C. One aliquot was used to determine the TCID₅₀ and the focus forming unit (FFU) titers of the stock.

Antiviral screening of compounds for activity against FeLV-FAIDS was performed in 96-well trays (Corning). This screening procedure is a modification of the FeLV infectivity assay established by Fischinger et al., (1974). Forty-eight hours prior to the assay, the indicator cells, 81C (obtained from D. Graves, University of Oklahoma, Oklahoma City, OK) were subcultured at a ratio of 1:2. Twenty hours prior to the assay, 96-wells trays were seeded with the 81C cells at 5 x 10₃ cells/well. On the day of the assay, the cells were pretreated for 30 minutes at 37°C with DEAE-dextran (25 µg/ml) in 0.1 ml Hanks balanced salt solution. This was removed and then 0.1 ml of growth medium containing 32 TCID₅₀ of FeLV-FAIDS, or 0.1 ml of growth medium alone, was added to each well. The virus was allowed to adsorb for 1 hour, then 0.1 ml of test or control compound (2',3'-dideoxycytidine or 3'-azidothymidine), or growth medium was added. Plates were incubated at 37°C. Cells were fed fresh growth medium containing compound on day 4 post-infection. Culture media were completely changed and replaced with fresh media containing compound on day 7 post-infection. On day 10 post-infection the cells were fixed by addition of 100µl buffered formalin for at least 1 hour at room temperature all

liquid was then removed and then the cells were stained with 0.1% Coomassie Brilliant Blue R-250. The plates were rinsed with deionized water, allowed to dry, and then observed microscopically for CPE and drug cytotoxicity.

3.3.11.2.2 Simian Retrovirus - SAIDS (SRV-2)

We received from Dr. Che-Chung Tsai one T25 flask and one 15 ml tube of a co-culture of Raji cells infected with a simian type D retrovirus (SRV-2) isolated from a macaque with SAIDS at the Washington Regional Primate Research Center. Subsequently, we purchased uninfected Raji cells (CCL-86) from the ATCC for the purpose of making fresh co-cultures with which to propagate the virus. These cells were successfully gown in Iscove's Modified Dulbecco's Medium containing 15% fetal bovine serum (complete Iscove's). The cells were subcultured every three days by a split ratio of 1:3. This yielded approximately 3 - 5 x 10⁵ cells/ml.

Co-cultures of normal Raji cells and Raji cells infected with SRV-2 were also maintained in complete Iscove's. These cells were subcultured every three days by a split ratio of 1:3 (yield = $3 - 5 \times 10^5$ cells/ml). Raji cells and infected co-cultures were maintained regularly. Fresh co-cultures were routinely set up according to the procedure provided by Dr. Che-Chung Tsai of the Washington Regional Primate Research Center. Briefly, 1 ml of 8 ml of an actively growing co-culture was added to a T25 flask containing 8 ml of complete Iscove's medium to which 0.5 ml of actively growing Raji cells had been added. This new co-culture was incubated at 37° C with 5% CO₂ with the flask in an upright position.

The antiviral screening of compounds against the SAIDS virus was performed by a syncytia-inhibition assay on Raji cells. Compounds were diluted in complete Iscove's medium and then $100~\mu l$ of each dilution was added to the appropriate wells of a 96-well plate. Actively growing Raji cells, 5~x 10^3 cells in $50~\mu l$ of complete Iscove's medium, were then added to each well. This was followed by the addition of $50~\mu l$ of clarified supernate from an SRV-2/Raji cell co-culture. Plates were incubated at $37^{\circ}C$ in a humidified atmosphere containing 5% CO_2 . Syncytia were counted on day seven post-infection. Drug toxicities were ascertained by comparing viable cell counts of uninfected, drug-treated samples to the viabilities of uninfected, untreated controls.

3.3.11.2.3 **Murine AIDS (MAIDS)**

Drug testing was begun in the CAS-BR-M murine leukemia virus plaque-reduction assay on compounds which had demonstrated significant antiviral activity in the HIV primary screen. The CAS-BR-M murine leukemia virus was obtained from Dr. John Billelo, V.A. Medical Center, Baltimore, MD. Dr. Billelo provided a culture of infected SC-1 producer cells which were propagated, and the supernatant containing infectious virus was used for the stock virus. These producer cells were maintained in Dulbecco's MEM supplemented with 10% heat-inactivated fetal bovine serum.

The continuous passage SC-1 feral mouse embryo cell line and the continuous passage rat XC sarcoma cell line (obtained from the American Type Culture Collection, Rockville, MD) were used for the plaque assay. Eagle's MEM with Earle's balance salt solution supplemented with 5% or 10% heatinactivated FBS was used as the growth medium for the propagation of both cell lines.

Falcon 6-well tissue culture plates were seeded with 1.75×10^5 cells per well in a total volume of 2.5 ml EMEM containing 5% inactivated FBS. Twenty hours after the cells were seeded, the medium was decanted and 2.5 ml DEAE-dextran (25 μ g/ml in phosphate buffered saline) was added to each well. The cultures were incubated at 37°C for 1 hour, after which the DEAE-dextran solution was decanted and the cell layers rinsed once with 2.5 ml PBS. Normal cell controls were refed with 2.5 ml medium alone (no virus or drug). Drug control cultures received 2.5 ml of medium containing drug but no virus.

Virus-infected control cultures received 0.5 ml of the appropriate dilution of stock CAS-BR-M to produce countable plaques plus 2.0 ml medium. The test samples received 0.5 ml of the appropriate virus dilution plus 2.0 ml of the appropriate drug dilution. Six concentrations of each test compound diluted in serial half-log₁₀ dilutions were tested.

Triplicate wells for each concentration of test compound and 6 virus and 6 cell control cultures were included in each assay. Hemacytometer cell counts, using the trypan blue dye-exclusion test for viability were done on duplicate cell control cultures at the time of virus inoculation.

On day three post-virus inoculation, the cultures were irradiated with an ultraviolet lamp for 20 seconds and XC cells were added to each culture (5 x 10⁵ cells/well in 2.5 ml EMEM containing 10% inactivated FBS). Viable cell counts were made on control and drug-treated SC-1 cell cultures.

On day three post-UV irradiation, the cultures were fixed with formalin and stained with crystal violet. The plaques were counted with the aid of a dissection microscope.

Antiviral activity in the CAS-BR-M plaque-reduction assay was expressed in terms of the reduction in the mean number of plaques counted in the drug-treated, virus-infected cultures compared with the mean number of plaques counted in the untreated, virus-infected control cultures (percent of control). The MIC₅₀ and a TI were also calculated.

3.4 Prescreen Assays (YF, VE, PT):

Large numbers of plant extracts are available for screening for in vitro antiviral activity. We were requested to develop an assay that would: 1) allow more compounds to be evaluated per microtiter plate than in the regular antiviral screen and 2) prescreen plant extracts as well as other compounds for activity against two indicator viruses. The three viruses selected by USAMRIID for this purpose were the attenuated, vaccine strain (17D) of Yellow Fever virus (YF) and Punta Toro virus (PT) and Venezuelan Equine Encephalomyelitis virus (VE). The prescreen assay should select candidate compounds for screening against the Asibi strain of YF and the other target RNA viruses in the full screen.

A prescreen procedure was developed which utilizes MTT and evaluates five compounds per virus per 96-well plate.

Host Cells. Vero cells are seeded as monolayer cultures in COSTAR 96-well plates at 18,000/0.2 ml/well in MEM + 10% heat-inactivated fetal bovine serum (Δ fbs). The plates are incubated approximately 24 hours prior to use.

Challenge Virus. The 17D strain of YF virus was originally obtained by Dr. Lori Brando from the Alabama Public Health Department in Birmingham, Alabama. The Adames strain of PT virus was obtained from our regular screen. Stock virus pools were prepared by passaging and titrating each virus in Vero cell monolayers. For use in the prescreen assays, PT virus is diluted in experiment medium (MEM + 2% Δ fbs) to yield an inoculum of 32 CCID₅₀ per well. The challenge dose of YF virus had to be increased from 32 to 64 CCID₅₀/well in order to obtain sufficient CPE and cell-kill by 6 days post-infection. An antiviral screening assay as long as 7 or 8 days would require replacing the culture fluids with fresh drug during the incubation period, thus increasing the cost of the assay.

Test compounds are dissolved or suspended in DMSO or H_2O , then diluted in serial tenfold dilutions in experiment medium to yield final concentrations of 1000, 100, 10 and 1 μ g/ml in the plate wells. Selenazofurin (AVS-0253) is utilized as a positive control drug for YF and VE; the control drug for PT is Ribavirin (AVS-0001). AVS-6724 (2-Thio-6-Azauridine) has been tested against all three viruses as a possible candidate positive control drug.

Assay Procedure. The assay is designed to evaluate four 1.0 \log_{10} concentrations of each compound in duplicate against the challenge virus. MEM + 2% Δ fbs serves as the experiment medium. Cell culture medium is removed from the plate wells. To each of duplicate test wells containing replicate cell monolayers, 100 μ l of each test drug solution (or suspension) and 100 μ l of the challenge virus (diluted in experiment medium) are dispensed. Cell controls (6/plate) containing 200 μ l medium, virus-infected, untreated cell controls (6/plate) containing virus and medium and drug cytotoxicity controls (1/drug concentration) containing cells, medium and test drug are included on each plate. Medium (Reagent) color controls (no cells), and drug colorimetric controls (drug + medium - no cells) accompany each test. The covered plates are incubated at 37°C in a humidified atmosphere containing 2% CO₂. When CPE reach 100%, 6 days post infection, 20 μ l aliquots of MTT (5 μ g/ml solution in PBS) are added to each well. The plates are incubated at 37°C for six hours to allow reduction of the MTT to the formazan form. Then 40 μ l aliquots of a 30% solution of SDS in 0.02 NHCl are added to the plate wells. The plates are incubated overnight to allow the SDS to lyse the cells and to dissolve the

MTT formazan crystals. The absorbance (570 nm) of the contents of each well is determined by a plate reader. The plate reader is interfaced with a computer programmed to capture the optical density (O.D.) measurements from the reader and calculate mean optical densities, indices such as IC_{50} , TC_{25} , SI, and plot the percents cell viability and CPE reduction. The actual O.D. readings, the indices, and bar graphs expressing cell viability and CPE reduction are automatically printed on individual compound data sheets.

An example of a data sheet with results of a prescreen assay is shown as Figure 2. Section I of the data sheet gives the compound and test identification and the actual raw data (O.D. readings) for each assay. Printouts of pertinent Control and Test Results are in Section II. Calculations of the test results are the same as those for assays in the regular screen. Section III displays a bar graph plotted from the computed values in Section II.

IN VITRO ANTIVIRAL RESULTS

EXOTIC RNA VIRUSES PRE-SCREEN ASSAY

DRUG: AVS 0001

SI: >5.04 08/24/89 PLATE NUMBER TEST DATE 08/31/89 PROJECT 5975-4 DATE READ SPONSOR USAMRIID CELLS VIRUS Satisfactory STRAIN ADAMES 10 11 12 drug A - 0001 Section I 1.554 1.537 0.155 1.471 3 0.510 0.686 0.155 1.537 C 0.231 0.206 0.178 1.506 0.223 0.227 D 0.157 virue control reagent E 0.211 0.361 0.220 0.175 P 10.249 0.278 0.325 0.179 G 11.455 1.530 0.174 1.493 (1.402 1.474 0.177 cell control blank BOLD - highest drug conc

AVS 0001 vs. PT (08/24/89)

UNINFECTED INFECTED REAGENT 0.176 ROW ON . COMC. CELL VIABILITY CPE REDUCTION VIRUS CONTROL 0.098 PERCENT (uG/mL) MEAN PERCENT CELL CONTROL 1.300 MEAN 100% 1.247 1009 DIFFERENTIAL 1.202 100 1.399 Section II A B 10 1.316 100% 0.346 29% IC50 19.80 C 1 1:359 100% -0.057 0 TC25 100.00 0.1 1.349 100% -0.030 0 SI 5.04 100% 90% (%) 80% CONTROL CELL VABILITY 70% 60% Section III 50% 40% 8 30% 20% 10% CC VC 100 DRUG CONCENTRATION (ug/mL) INFECTED CELLS UNINFECTED CELLS Prescreen Data Sheet Figure 2

3.5 Antiviral Evaluation In Vivo:

3.5.1. Pichinde Virus in MHA Hamsters (PIC):

The in vivo testing protocol used for evaluation of antiviral efficacy against Pichinde virus was as follows: Soluble compounds were diluted in phosphate-buffered saline (PBS) (pH 7.2) while insoluble compounds were diluted and homogenized in 0.4% carboxymethylcellulose (CMC) in PBS. Other diluents were used specified by the sponsor. If the test compounds were stable, then compound solutions (suspensions) were prepared for the first half of the experiment on the first day of dosing. The solutions/suspensions for the second half of the experiment were prepared on Day 3. The compound solutions/suspensions were held at 4°C when not in use. Compounds with known instabilities were prepare daily prior to administration. In general, the compounds were prepared at concentrations appropriate for dosing hamsters at 0.1 ml of solution per 20 gm of body weight. Fifteen MHA strain hamsters (4 wk old, 60-80 gm) (5 uninfected, 10 infected) were treated subcutaneously at each dose level once daily for 7 days (qld x 7) with the first dose given 2 hr preceding or 4 hr following virus challenge. The virus challenge consisted of 1 x 10⁴ pfu of plaque-purified Pichinde virus administered intraperitoneally in a volume of 0.5 ml. Compounds were administered to each hamster based on the average cage weight (5 hamsters/cage). Treatment groups included the test compounds as well as the diluent used for compound preparation. Controls included (1) untreated, uninfected hamsters, (2) untreated, virus-infected hamsters, (3) virus-infected diluent-treated hamsters and (4) ribavirin (positive control drug) treated hamsters.

Hamsters were monitored daily for mortality and body weight changes through day 28 post-infection. The mortality rates occurring in drug-treated hamsters were compared using Fisher's Exact test. The average day of death (ADD) was calculated and compared by Student's t-test. The virus rating (VR) was calculated as the geometric mean time to death (GMTD) of the drug treated hamsters divided by the GMTD of the diluent treated controls.

In several instances, this protocol was modified so that the compound were administered on alternate treatment schedules (i.e. qld x 10 days).

3.5.2. Venezuelan Equine Encephalomyelitis Virus (VE):

A prophylactic treatment protocol was used for evaluating the antiviral activity of compounds against Venezuelan Equine Encephalomyelitis virus infection of CD-1[®] outbred mice. Compounds submitted for in vivo evaluation were diluted in PBS (pH 7.2) if they were soluble or they were diluted and homogenized in 0.4% CMC in PBS if they were insoluble. Compounds were prepared at concentrations consistent with a dose volume of 0.1 ml of compound per 10 gm of body weight. Adequate compound was prepared for the entire treatment schedule unless the compound data sheet indicated that the compound was unstable. Unstable compounds were prepared immediately before each administration. Diluted compounds were held at 4°C when not in use. All compound administration was by the SC route on an individual animal weight basis. Fifteen female CD-1° mice (18-21 gm) were treated at each compound dose level of a qld x 7 days schedule starting day -1. Ten of the 15 mice were challenged IP with an LD₉₀ dose of virus in a volume of 0.5 ml of MEM/2% FCS. The remaining 5 mice served as toxicity controls. Because of compound quantity limitations, the highest dose of compound generally tested was 320 mg/kg with 4 serial half-log dilutions below this level. Treatment groups included the test compound dilutions as well as the diluent used for compound preparation. Controls included (1) untreated, uninfected animals, (2) untreated, infected animals, and (3) virus-diluent treated animals (infected and uninfected). No positive control drug was available for this model. Mortality was monitored for 21 days post-infection. The calculations and statistical comparisons were as described for Pichinde virus.

3.5.3. Japanese Encephalitis Virus:

A prophylactic treatment protocol was used for evaluating the antiviral activity of compounds against Japanese Encephalitis virus infection of inbred C57Bl/6 mice. Compounds submitted for in vivo evaluation were diluted in PBS (pH 7.2) if they were soluble or they were diluted and homogenized in 0.4% CMC in PBS if they were insoluble. In some cases, compounds were solubilized in ethanol or DMSO prior to dilution into PBS or 0.4% CMC/PBS. Compounds were prepared at concentrations consistent with a dose volume of 0.1 ml of compound per 10 gm of body weight. If there were no stability problems noted on the compound data sheet then compound quantities adequate for the first half of the experiment were prepared on the first day of dosing. The remaining compound was prepared on experimental day 2. Unstable compounds were prepared immediately before administration. Diluted compounds were held at 4°C when not in use. Compound administration was primarily by the SC route on an individual animal weight basis; however, in some cases the IP route was used for comparative purposes. Fifteen female C57Bl/6 mice were treated at each compound dose level on a gld x 7 days schedule starting day -1. Ten of the 15 mice were challenged IP with an LD₉₀ dose of virus in a volume of 0.5 ml of MEM/2% FCS. The remaining 5 mice served as toxicity controls. Because of compound quantity limitations, the highest dose of compound generally tested was 320 mg/kg with 4 serial half-log dilutions below this level. Treatment groups included the test compound dilutions as well as the diluent used for compound preparation. Controls included (1) untreated, uninfected animals, (2) untreated, infected animals and (3) virus diluent treated animals (infected and uninfected). No positive control drug was available for this model. The mice were monitored for 21 days. Calculations and statistics were as described for Pichinde virus.

3.5.4. Vaccinia Virus:

3.5.4.1. Intracranial Challenge:

A prophylactic treatment protocol was used for evaluating the antiviral activity of compounds against Vaccinia virus infection of CD-1[®] outbred mice. Compounds submitted for in vivo evaluation were diluted in PBS (pH 7.2) if they were soluble or they were diluted and homogenized in 0.4% CMC in PBS if they were insoluble. In some instances the compounds were solubilized in DMSO prior to dilution in PBS or 0.4% CMC/PBS. Compounds were prepared at concentrations consistent with a dose volume of 0.1 ml of compound per 10 gm of body weight. If there were no stability problems noted on the compound data sheet then compound quantities adequate for the first half of the experiment were prepared on the first day of dosing. The remaining compound was prepared on experiment day 2. Unstable compounds were prepared immediately before administration. Diluted compounds were held at 4°C when not in use. Compound administration was by the SC or by the IP route on an individual animal weight basis. Fifteen female CD-1 mice (18-21 gm) were treated at each compound dose level on a qld x 7 days schedule starting day -1. Ten of the 15 mice were challenged intracranially with an LD₀₀ dose of virus in a volume of 0.03 ml of MEM/2% FCS. The remaining 5 mice served as toxicity control. Because of compound quantity limitations, the highest dose of compound generally tested was 320 mg/kg with 4 serial half-log dilutions below this level. Treatment groups included the test compound dilutions as well as the diluent used for compound preparation. Controls included (1) untreated, uninfected animals (2) untreated, infected animals and (3) virus diluent treated animals. Ara-A was the positive control drug for this model.

3.5.4.2. Tailpox Model:

The IHD strain of vaccinia virus, passed once in mouse brain and once in primary rabbit kidney cell culture, was used. Random-bred Swiss mice (CD-1TM, VAF+, Charles River Laboratories, Inc.), weighing 18-21 grams, were inoculated via the tail vein (1 cm from the base) with 0.2 ml of a 1:40 dilution of the virus suspension. Compounds were administered SC once daily for 7 days with the first

dose given the day preceding virus challenge. Uninfected drug-treated toxicity controls were included for each treatment administered. Animals were sacrificed on the sixth day and their tails were stained with a solution of 1% fluorescein-0.5% methylene blue in 70% methanol. Lesions were enumerated under ultraviolet light (354 nm) with the aid of a hand lens. The average number of lesions for each treatment group was calculated prior to, and following square root transformation of the individual tailpox counts. Tailpox counts from each of the treatment groups were compared by student t-test. A p-value at or below 0.05 was considered indicative of antiviral activity.

4. RESULTS

4.1 In Vitro Antiviral Evaluations: DNA, Exotic RNA and Retrovirus Viruses:

During this five-year reporting period from November 16, 1985 to January 31, 1991, a total of 9658 (5849 primary screen and 3809 prescreen) test compounds were received at SRI for evaluation in the *in vitro* antiviral screen (prescreen and primary). A cumulative summary of the compound shipments received, and the number of compounds in each shipment, are presented in Table 2. Compounds received in amounts too small for appropriate evaluation against all of the viruses were screened according to a priority list (determined by the sponsor) of the target viruses from the primary screen. In some instances, the testing of compounds against the exotic viruses was coordinated with the testing against Human Immunodeficiency Virus (HIV). *In vitro* primary screening evaluations were carried out against Vaccinia Virus (VV), Adenovirus Type 2 (AD2), Vesicular Stomatitis Virus (VSV), Yellow Fever Virus (YF), Japanese Encephalitis Virus (JE), Venezuelan Equine Encephalomyelitis Virus (VE), Punta Toro Virus (PT), Sandfly Virus (SF), Pichinde Virus (PIC), and Hantaan Virus (HTN). In July 1986, the HIV virus was added as an additional virus to be tested under the primary screen. In January, 1988, primary screening of compounds versus HIV was moved from this contract to a cooperative agreement identified as DAMD17-88-H-8003 with USAMRIID.

Approximately 66,000 in vitro antiviral assays were performed during this contract period. This number includes all quality control and unsatisfactory tests (Figure 3). Positive control drugs, as specified in Table 1 were tested in parallel in each assay. Numerous internal virus load and cell load quality control tests were performed. Results of the cell controls and virus controls were monitored to test for viability, consistency and repeatable results during the day-to-day operation. Figure 4 illustrates the number of compounds that were tested in vitro primary screen each year.

The results for compounds found active are summarized in the following sections for each virus. A cumulative summary presenting all of the *in vitro* antiviral tests results (both positive and negative) is included in Appendix A and B.

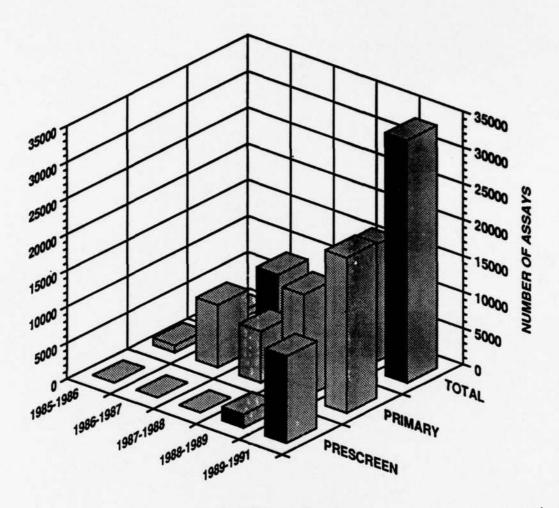
Primary Testing of compounds versus HIV was performed for the USAMRIID program by using the modified Broder assay. In order to accommodate the large volume of HIV screening required by the NCI modification to this contract, testing was carried out using the XTT automated assay system.

Table 2

Prescreen and Primary In Vitro Antiviral Screen Number of Compounds Received Per Shipment Through January 31, 1991

Shipment	Per	Date	Shipment	Per	
Number	Shipment	Received	Number	Shipment	Received
34	49	11/25/87	3	220	04/10/90
35	8	12/22/87	19	95	05/00/50
36	65	01/29/88	89	201	05/30/90
37	84	03/09/88	69	191	06/27/90
38	S	. 38/21/80	02	156	06/53/90
39	89	03/23/88	. 71	S	07/18/90
40	57	04/26/88	72	122	04/18/90
41	99	05/13/88	73	142	09/11/90
41B	93	05/13/88	74	991	10/04/90
41C	4	05/13/88	75	187	10/25/90
42	%	05/13/88	92	320	12/06/90
43	140	07/08/88			
\$	68	08/10/88	SubTotal		5849
45	9/	09/13/88	Prescreen		
46	126	10/06/88	IP	869	68/80/90
47	7	10/20/88	2P	12	07/20/89
48	101	11/02/88	3P	695	08/56/89
49	1	11/11/88	4P	38	02/01/90
20	1	11/16/88	4P	707	03/15/90
51	%	01/10/89	SP	168	06/01/90
52	8	05/09/89	6P	183	04/02/00
53	66	03/15/89	7P	152	08/22/90
54	71	04/19/89	8P	111	08/28/90
26	103	05/12/89	В	200	09/25/90
57	133	68/L0/90	10P	200	10/15/90
58	73	07/13/89	11P	200	11/16/90
59	49	08/11/80	12P	189	12/03/90
09	78	68/L0/60	13P	191	12/17/90
61	138	10/06/89	14P	195	1/24/90
62	991	11/21/89			
63	135	05/01/90	SubTotal		3809
2	4	05/27/90			
99	125	03/16/90	Total		9658

TOTAL NUMBER OF IN VITRO ANTIVIRAL ASSAYS (PRESCREEN AND PRIMARY SYSTEMS)

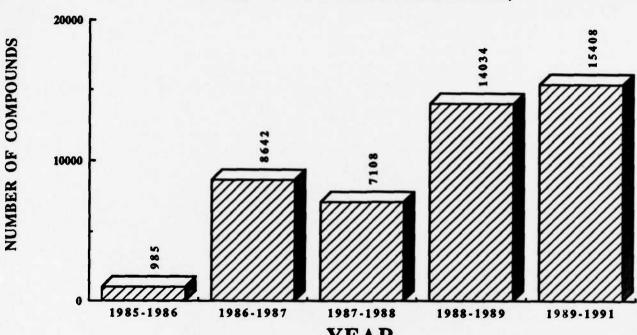


Number of Assays Tested

Year	Primary- Screen	Pre- Screen	Yearly <u>Total</u>
1985 - 1986	985	0	985
1986 - 1987	8,642	0	8,642
1987 - 1988	7,108	0	7,108
1988 - 1989	14,290	1,878	16,168
1989 - 1991	21,470	12,050	33,520
Total per Screen	52,495	13,928	Grand Tot 66,423

Figure 3

NUMBER OF COMPOUNDS TESTED IN VITRO PER YEAR DURING THE CONTRACT PERIOD (INCLUDES ALL VIRUSES TESTED IN THE PRIMARY SCREEN)



¥?			YEAR			Total No. Compounds Per Virus
Virus			***	626	1620	
VV	231	669	669	575	1520	3664
AD2	231	668	656	523	Not Required	2078
YF	0	819	875	1841	2717	6252
JE	0	905	848	1847	2710	6310
VE	0	167	854	2331	2715	6067
PT	138	871	869	1751	3005	6634
SF	139	718	877	1873	2741	6348
HTN	0	•	35			35
PIC	13	114	819	1195	•	2141
VSV	233	627	605	665	Not Required	2130
HIV (USAMRIID)	0	495	**	**	**	495
HIV (NCI)	0	1688	••	••	••	1688
TOTAL	985	7972	7107	12601	15408	44073
* Testing moved to			RIID).			(Grand Total) 44,073

^{*} Testing moved to Fort Detrick Campus (USAMRIID).

Figure 4

^{**} Testing transferred to another project.

4.1.1 Vaccinia Virus (VV):

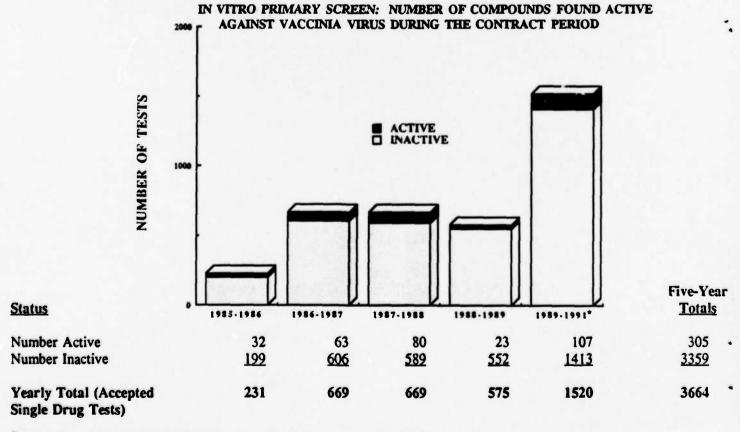
The number of single drug tests carried out against VV during this contract period is summarized in yearly increments in Figure 5. During this five-year period two main in vitro antiviral assay protocols were implemented:

- 1. A standard CPE inhibition assay by virus rating (VR) (Annual Report, December 15, 1988, Section 3.2.4).
- 2. Since January, 1989, MTT based-antiviral assay format.

A total of 4354 tests were performed during this contract period using both assay types. Routine testing was changed to the MTT-assay format to improve the efficiency and quality of the primary screening program in addition to being more cost-effective. Ara-A (AVS-1752) and Selenazofurin (AVS-0253) were tested in each standard virus rating (VR) CPE-inhibition assay as positive control compounds. Results of these positive controls (VR tests) were used as a guideline to assess the quality of each assay.

After the testing was converted to the MTT-assay format, we performed a total of 219 control compound assays with Ara-A and Selenazofurin during the last 24 months of the contract period. The rest, totaling 2095, were actual single drug MTT-assays. The total number of MTT-assays (2554) tested during the last two years represents a 42% increase (improvement) in the total testing output as compared to the total of 1800 tests performed during the first 3 years of this contract.

Out of the 3664 accepted single drug tests, 305 compounds demonstrated antiviral activity at greater than 50% reduction levels. This represents around 8.3% of the tested compounds having *in vitro* antiviral activity against VV-virus. The remainder, 3359 compounds (91.7%), were considered inactive with both assay protocols (Figure 5).



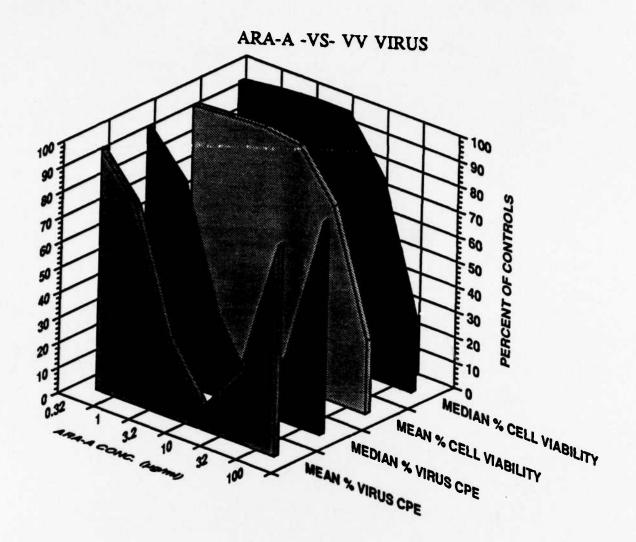
Represents 14-month period (November 15, 1989 - January 31, 1991)

4.1.1.1 <u>VV-Quality Controls (MTT Assay)</u>: Two positive control compounds (Ara-A and Selenazofurin) were used in the daily assay sets as antiviral activity quality controls. The antiviral performance of the unknown compounds is compared to that of the positive control compounds. Compounds with equal to or better antiviral potency are considered active and are worthy of further in vitro profile studies and in vivo testing.

The following indices summarize the Ara-A data.

	IC ₅₀ μg/ml	TC ₅₀ µg/ml	<u>SI</u>	TAI
Mean	2.17	69.26	18.52	42.81
S.D.	0.79	15.57	7.66	8.59
Median	2.02	71.9	16.38	42.76

The closeness of the mean and median values indicate the reproducibility of the effect of Ara-A on VV and on host cells. A summary of the antiviral and cytotoxicity values for 89 Positive Control Tests at each concentration of Ara-A are listed and plotted in Figure 6-A. Excellent reproducibility was achieved in the 89 tests performed during November, 1989 through January, 1991. Ara-A continues to be a reliable positive control drug.



CONCENTRATION (µg/ml)

		9	Viral	CPE					% Cel	l Viabil	ity	
Conc.(µg/ml)	0.32	1.0	3.2	10	32	100	0.32	1.0	3.2	10	32	100
Mean	95	78	31	9	29	84	96	97	98	96	77	29
Median	95	79	31	7	26	85	97	99	100	100	79	29
Std. Dev.	0.04	0.10	0.15	0.09	0.14	0.09	0.04	0.04	0.03	0.05	0.13	0.09

Figure 6-A

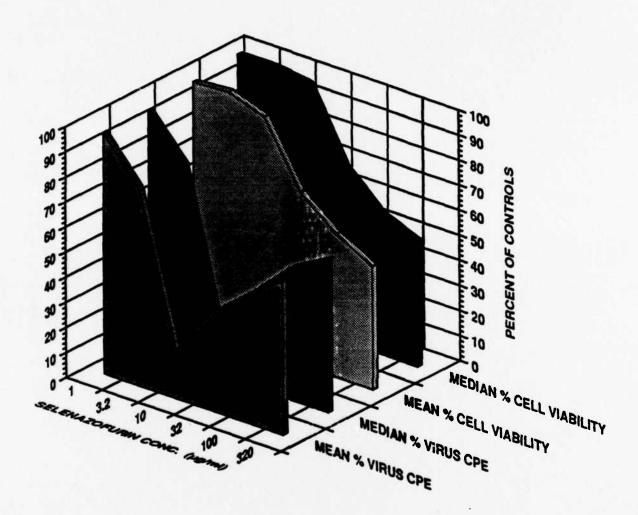
Average Antiviral and Cytotoxicity Values for 89 Positive Control Compound Tests

The following indices summarize the data for Selenazofurin:

	IC ₅₀ µg/ml	TC ₅₀ µg/ml	SI	TAI
Mean	5.70	233	6.51	27.61
S.D.	1.80	106	7.87	8.61
Median	5.40	320	4.80	26.52

As can be seen from the S.D.'s, cytotoxicity (TC₅₀) of Selenazofurin was somewhat inconsistent against VV in Vero cell cultures. This inconsistency is not significant since we are testing at 0.5 log₁₀ dilutions. However, if one examines the average antiviral and cytotoxicity values at each concentration of Selenazofurin from 83 assays that were performed during November, 1989 through January, 1991 (Figure 6-B), the results were quite reproducible.

SELENAZOFURIN -VS- VV VIRUS



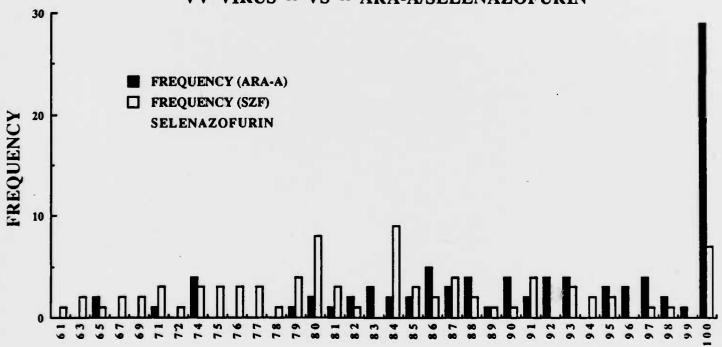
CONCENTRATION (µg/ml)

		90	Viral	CPE					% Cel	l Viabil	ity	
Couc.(µg/ml)	1	3.2	10	32	100	320	1	3.2	10	32	100	320
Mean	96	78	19	39	51	61	98	99	94	69	57	49
Median	96	79	16	39	53	61	100	100	98	70	56	50
Std. Dev.	0.05	0.11	0.12	0.11	0.10	0.11	0.03	0.02	0.08	0.09	0.09	0.10

Figure 6-B
Average Antiviral and Cytotoxicity Values for 83 Positive Control Compound Tests

4.1.1.2 <u>Maximum Antiviral Effect of Ara-A and Selenazofurin vs VV Virus:</u> The distribution of the maximum VV CPE inhibition values (% CPE reduction) recorded from the 172 positive control assays (89 tests were Ara-A and 83 tests were Selenazofurin) conducted during the period of November, 1989 through January, 1991, is depicted in Figure 7. As expected, Ara-A had a much greater inhibitory effect than Selenazofurin.

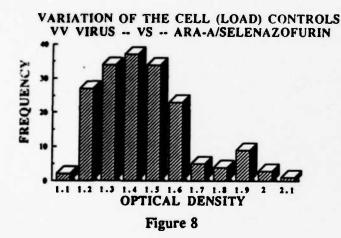
VARIATION OF THE MAXIMUM ANTIVIRAL EFFECT VV VIRUS -- VS -- ARA-A/SELENAZOFURIN

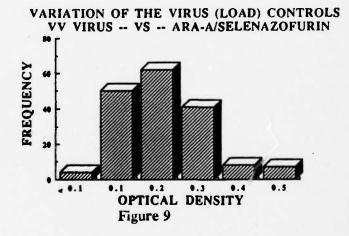


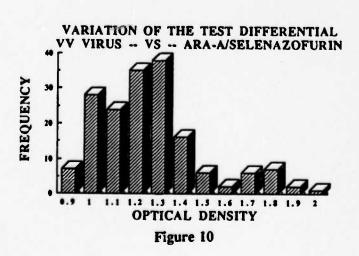
PERCENT CPE REDUCTION

Figure 7

4.1.1.3 <u>VV-Assay Plate Quality Controls: Cell Load and Virus Load Parameters (Ara-A/Selenazofurin):</u> The MTT assay is fundamentally dependent upon the quality of the assay plates. Our large-scale antiviral testing is dependent upon the uniformity of the test plates produced for the daily assays. Equal numbers of cell load and virus load as well as the consistent performance of the reagents used daily was monitored. A sample of the plate variation control for the period of November, 1989 through January, 1991, is presented in Figures 8, 9 and 10. The cell control data (Figure 8) indicates that the cell load was somewhat less consistent than the virus load during the past fourteen-months. The plot of the virus control O.D.'s (Figure 9) indicates good host cell destruction and a uniform load of virus among the day-to-day experiments. Figure 10 gives the frequency and variation of the mean values (O.D.) of the differential from each positive control assay. Each differential value represents the difference between the mean cell control O.D. and the mean O.D. of the virus controls of each of the 172 assays.







4.1.1.4 <u>VV-Antiviral Activity Results:</u>

New Drugs with 95% Antiviral Reduction Levels: Out of the 3895 actual single drug tests, 34 new compounds demonstrated excellent antiviral activity, having antiviral reduction values of equal to or better than 95% and Selectivity Indices (SI) of ≥ 1.0 . This represents around 0.9% of the test compounds being active at this maximum reduction level criteria. These compounds are summarized in **Table 3** according to the highest **Total Antiviral Index (TAI)**. Compounds AVS-5569, 5568, 1214, 5219, 0303, 7449 and 6193 demonstrated the greatest *in vitro* promise, having TAI's that ranged from 51 - 83 and SI's that ranged from 19 - > 320. The next thirteen compounds, AVS-3571, 1654, 8586, 6218, 2893, 6215, 5035, 6210, 2568, 6462, 2275, 1211 and 6220, demonstrated excellent antiviral activity with TAI's that ranged from 30 - 48 and SI values that ranged from 6 - 21. The rest (12 compounds) had only moderate antiviral activity with TAI's ranging from 11 - 27 and SI's of 1 to 5.9.

It is worthwhile to note that compounds received in shipment number 62 were mostly colored (Table 3). Therefore those compounds appearing in the 95% active category from shipment number 62 should be interpreted with caution, since colored compounds can create false positive readings with the MTT assay. However, the drug color controls (OD's) are subtracted from the Test OD's and should compensate for higher readings due to drug color.

Table 3 ${\tt AVS~Compounds~Active~Against~Vaccinia~Virus~(VV)~at~{\tt AI}_{95}~{\tt Level}}$

AVS	Ship-		Diff-					
No.	ment#	Date	rnt1.	IC 95	TC 95	AI 95	SI	TAI
** VIRUS								
VV 5569		07/13/89	1.469	2.17 >	320.00 >	147.36 >		
VV 5568	09	07/13/89		2.17 >	320.00 >	147.36 >		
VV 1214	52	03/23/89		12.10 >	320.00 >	26.40		> 65.14
VV 5219 VV 0303	51 62	05/04/89		0.79 > 0.92 >	10.00 >	12.70		> 62.29
VV 7449	71	01/17/90 10/04/90		0.92 >	32.00 > 100.00 >	34.66		> 59.69 > 59.01
VV 6193	62	02/22/90		191.00 >	320.00 >	114.81 >		> 51.01
VV 3571	32	08/17/89		30.90 >	320.00 >	10.36 >		> 47.60
VV 1654		06/01/89		0.96 >	32.00 >	33.40		> 46.18
VV 8586	76	01/31/91		100.00 >	320.00 >	3.20	20.95	39.42
VV 6218	62	02/22/90		277.00 >	320.00 >	1.16 >		> 37.51
VV 2893	26	08/17/89		78.90 >	320.00 >	4.06	10.26	37.30
VV 6215	62	01/24/90	1.209	91.60 >	320.00 >	3.49 >	7.69	> 36.32
VV 5035	48	06/01/89	2.287	6.66 >	32.00 >	4.81	8.97	34.26
VV 6210	62	02/22/90		93.00 >	320.00 >	3.44 >		> 33.52
VV 2568	67	06/27/90		29.60 >	100.00 >	3.38 >		> 33.06
VV 6462	63	03/29/90		9.54 >	320.00 >	33.56		> 32.30
VV 2275	67	06/21/90		29.30 >	320.00 >	10.91	10.01	32.18
VV 1211	27	08/17/89		72.90 >	320.00 >	4.39	8.48	32.10
VV 5053 VV 6220	48 62	01/05/89		30.10	274.00 320.00 >	9.11 3.39 >	5.20	31.95
VV 5069	48	01/17/90 01/12/89		94.5C > 264.00 >	320.00 >	1.21 >		> 30.04 > 26.64
VV 0206	67	06/14/90		93.90 >	320.00 >	3.41		> 26.53
VV 7053	72	10/04/90	1.113	282.00 >	320.00 >	1.13 >		> 26.45
VV 6209	62	03/15/90		89.60 >	100.00 >	1.12 >		> 24.55
VV 6219	62	02/22/90		264.00 >	320.00 >	1.21 >		> 24.05
VV 5072	48	09/07/89		139.00 >	320.00 >	2.30		> 22.92
VV 6986	68	07/19/90		92.70 >	320.00 >	3.45	3.89	21.69
VV 2866	20	08/17/89		289.00 >	320.00 >	1.11 >		> 20.04
VV 6199	62	01/11/90		92.80	309.00	3.33	3.28	19.71
VV 6217	62	04/12/90	1.265	92.20 >	100.00 >	1.08 >	2.26	> 19.24
VV 6192	62	02/22/90		94.50 >	100.00 >	1.06 >		> 14.61
VV 7479	73	11/29/90		315.00 >	320.00 >	1.02 >		> 13.67
VV 7023	69	07/12/90	1.299	301.00 >	320.00 >	1.06 >	1.84	> 11.28
DIFRNTL	=	The difference densities.	ential is the	e difference in	the cell con	trol and the	virus con	trol optical
IC ₉₅ =				on 95% = T by using a re				
$TC_{95} =$		toxicity con ty by 95%.	centration	95% = The	drug concent	ration (µg/	ml) that re	educed cell
$AI_{95} =$				point ration 6 5% reduction				
SI =				calculated by nl, the maxim			e IC ₅₀ (bas	sed upon 6
TAI =		Antiviral Ind scale of 0-		area between i	the cytotoxici	ty and the a	ntiviral cu	rves (hased

New Drugs with 50% Antiviral Reduction Levels: Out of the 3895 actual single drug tests, 91 new compounds demonstrated good antiviral activity, having antiviral reduction values equal to or better than 50% and SI's of > 1. This represents around 2.3% of the test compounds being active at this good antiviral reduction level criteria. These compounds are summarized in Table 4 according to the highest Total Antiviral Index (TAI). AVS-1985 and AVS-138 demonstrated the best TAI's of 73 and 63 and SI's of 178 and 103, respectively. Twenty-four other compounds demonstrated moderate antiviral activity, having TAI's that ranged from 20 - 53 and SI's from 2 - 32. The rest (65 compounds) showed marginal antiviral activity with TAI's that ranged from 3.4 to 19 and SI's from 1 to 5.5.

It is worthwhile to note (Table 4) that compounds received in shipment number 62 were mostly colored. Therefore those compounds appearing in the 50% active category from shipment number 62 should be interpreted with caution, since colored compounds create false positive readings with the MTT assay. However, the absorbance values of the drug color controls are subtracted from the test O.D.'s which should compensate for the increased absorbance due to drug color.

Table 4 ${\tt AVS~Compounds~Active~Against~Vaccinia~Virus~(VV)~at~{\tt AI}_{50}~{\tt Level}}$

AVS	Ship-	Test	Diff-							
No.	ment#		rnt1.		IC 50		TC 50	AI 50	SI	TAI
** VIRUS	VV									
VV 1985	67	06/21/90	1.077	•	0.03 >		10.00 >	312 50 >	177.92 >	73.32
VV 0138	٠,	08/17/89	1.236	•	3.11 >		320.00 >		102.95 >	
VV 7447	71	10/04/90			8.59 >		320.00 >	37.25	24.45 >	
VV 2290	71	08/02/90			8.75		286.00	32.69	21.84 >	
VV 1655	62	12/22/89			9.13		287.00	31.44	9.15 >	
VV 7448	71	10/04/90			32.00 >		320.00 >	9.99 >		
VV 0001	'-	06/14/90			27.10 >		320.00 >	11.81 >		
VV 5034	48	05/18/89			15.80 >		320.00 >	20.30 >		
VV 3802	67	06/27/90			0.10		20.80 >	208.25 >		
VV 5988	61	12/07/89		-	1.95		27.30	14.01	8.96 >	
VV 6724	67	06/27/90			3.81		85.80	22.51	7.28	30.67
VV 2860	20	09/07/89			6.10		63.20	10.36	6.60 >	_
VV 2318	67	07/19/90			0.15		19.20	123.29	11.58 >	
VV 5987	61	12/07/89			1.45		20.10	13.79	8.39 >	
VV 5450	53	04/13/89	1.537		6.20		98.10	15.80	8.21 >	
VV 6224	62	01/17/90			62.60 >	>	320.00 >	5.12 >	5.12 >	
VV 5997	66	06/14/90			0.63 >		100.00 >	159.38	10.16 >	
VV 2966	27	08/17/89			8.99		158.00	17.57	7.25 >	
VV 2503	67	07/19/90			0.17		8.57	49.93	7.96 >	
VV 6223	62	01/17/90			45.60		283.00	6.22	4.21	23.39
VV 6467	63	04/19/90			0.18		1.86	10.53	4.65	23.24
VV 0646	67	07/12/90			0.19		9.24	47.74	4.21	23.11
VV 8239	75	12/13/90			55.60		308.00	5.53	3.67	21.86
VV 6042	61	12/13/89			49.10		172.00	3.51	1.99 >	
VV 4934	51	02/02/89			11.20		50.50	4.53	1.68	20.86
VV 6469	63	03/29/90			43.20		251.00	5.81	3.91	19.64
VV 6304	63	03/01/90	1.154		67.30 >	>	320.00 >	4.76	4.63 >	19.07
VV 8532	76	01/24/91	1.224		13.80		73.10	5.30	2.73 >	19.00
VV 6754	67	07/19/90	1.305		1.38		8.93	6.45	3.21	18.78
VV 2991	41B	08/17/89	1.185		2.05		18.70	9.14	5.47	18.75
VV 5601	67	06/07/90			151.00 >	>	320.00 >	2.12 >		
VV 1644	64	04/12/90	1.163		58.60		293.00	5.00	3.35	17.96
VV 5958	60	12/07/89	1.407		5.91		37.90	6.41	3.63	17.94
VV 7469	73	11/01/90			88.80 >		320.00 >	3.60 >	3.60 >	
VV 0148	67	06/21/90			0.55 >	>	10.00 >	18.36	4.36	17.58
VV 8356	76	01/31/91			0.60		6.00	10.02	3.59 >	
VV 5186	58	09/07/89			49.80		299.00	6.01	2.56 >	
VV 7485	73	11/01/90			32.00		190.00	5.94	3.33	16.50
VV 6184	62	01/11/90			220.00 >	>	320.00 >	1.46 >	1.46 >	
VV 7413	70	09/20/90	1.209		19.60		86.60	4.43	3.00	16.22
VV 5457	54	06/08/89			5.70		20.90	3.66	2.60	15.75
VV 6973	68	06/27/90			133.00 >	•	320.00 >	2.40 >	2.40 >	
VV 6145	62	01/10/90			41.60		233.00	5.60	2.53	15.43
VV 7302	70	09/27/90			320.00 >		320.00 >	1.00 >	1.00 >	
VV 5069	48	04/06/89			86.90 >		320.00 >	3.68	3.39 >	
VV 6213	62	01/24/90			81.00 >	•	100.00 >	1.23	1.12 >	
VV 5977	61	03/29/90			0.57		2.29	4.00	2.12 >	
VV 1355	64	03/29/90	1.409		154.00 >		320.00 >	2.08 >		
VV 4855	48	12/29/88			9.00 >		10.00 >	1.10 >		
VV 5146 VV 5973	57	08/17/89			2.88 >		10.00 >	3.47 >		
VV 5973 VV 2743	61 41B	12/07/89			7.17		21.40	2.98	2.04 >	
VV 4532	41B 42	08/17/89 02/23/89			95.70 > 85.80 >		320.00 > 320.00 >	3.34	3.12 >	
VV 7110	70	08/09/90			42.90		191.00	3.73 > 4.45	3.73 > 2.56	
VV 8355	76	01/10/91			16.10		254.00			12.02
VV 4992	51	02/16/89			134.00 >		320.00 >	15.78	1.30	11.98
VV 0124	69	07/12/90	1 266		20.60	4	128.00	2.39 6.18	1.46 > 1.69	11.45
VV 4849	48	12/29/88			151.00 >		320.00 >	2.11 >	2.11 >	
** 1013	70	12/23/00	1.022		131.00 >		320.00 /	2.11 >	2.11 >	11.30

Table 4 (Cont'd)

	AVS	Ship-	Test	Diff-					
	No.	ment#	Date	rntl.	IC 50	TC 50	AI 50	SI	TAI
vv	1017	64	03/29/90	1.317	85.50	237.00	2.77	1.72 >	11.06
VV	2688	41B	08/17/89	1.325	3.05	9.38	3.08	2.07 >	10.95
VV	4531	42	08/17/89		135.00 >	320.00 >	2.38 >	2.38 >	10.89
VV	6133	62	01/04/90	1.310	21.50	52.10	2.42		10.87
VV	8357	76	01/10/91	1.091	302.00 >	320.00 >	1.06 >	1.06 >	
VV	6783	68	07/12/90	1.274	21.00	66.00	3.14		10.49
VV	7094	72	11/01/90	1.234	75.90	255.00	3.36	2.34	10.37
VV	5058	48	01/05/89	1.491	198.00 >	320.00 >	1.61 >	1.61 >	10.15
	5971	61	12/07/89	1.324	3.06	13.00	4.26	2.37	9.85
VV	7049	69	07/26/90	1.381	179.00 >	320.00 >	1.79 >	1.79 >	9.77
	7391	70	08/30/90	1.277	70.60	206.00	2.92		9.69
	6983	68	06/27/90	0.872	192.00 >	320.00 >	1.67 >	1.67 >	9.46
	5968	61	12/07/89	1.441	86.70	185.00	2.13	1.15	9.11
	4939	51	08/17/89	1.248	8.06	20.60	2.56	1.77	9.03
	6771	67	06/21/90	1.146	7.20	19.50	2.71	1.84	8.48
	4943	51	02/02/89	1.749	8.31	22.10	2.66	1.72	8.25
	7468	73	11/01/90	1.221	246.00 >	320.00 >	1.30 >	1.30 >	8.05
VV	7003	69	07/12/90	1.440	215.00 >	320.00 >	1.49	1.24 >	8.01
	6819	68	08/02/90	1.275	85.20	228.00	2.67	1.92 >	7.85
	7074	72	10/04/90	1.175	225.00 >	320.00 >	1.42 >	1.42 >	7.30
-	6212	62	03/15/90	1.105	71.40 >	100.00 >	1.40 >	1.40 >	6.95
	7022	69	07/12/90	1.436	69.40	152.00	2.18	1.16	6.74
-	6399	63	04/12/90	1.096	0.75	2.13	2.85	1.70	6.02
	5970	61	12/28/89	1.739	0.78	1.91	2.44	1.40	5.97
	6019	61	12/13/89	0.969	100.00	202.00	2.02	1.17	5.55
	7038	69	07/26/90	1.231	87.50	183.00	2.09	1.30	5.51
	1838	64	04/12/90	1.163	238.00 >	320.00 >	1.34	1.08 >	5.42
	7102	70	08/09/90	1.513	28.60	89.90	3.15	1.03	5.12
	5155	57	07/27/89	1.452	2.48	7.66	3.09	1.12	5.01
	5972	61	12/07/89	1.429	2.54	5.66	2.23	1.14 >	4.80
	5314	53	04/06/89	1.769	320.00 >	320.00 >	1.00	1.00 >	4.68
	6758	67	06/14/90	1.046	2.62	5.78	2.21	1.39	4.42
	5488	53	07/27/89	1.567	313.00 >	320.00 >	1.02 >	1.02 >	4.03
VV	4098	37	02/23/89	1.860	0.00 >	0.00 >	1.05 >	1.05 >	3.43

- DIFRNTL = The differential is the difference in the cell control and the virus control optical densities.
- IC_{50} = (Viral) inhibitory concentration 50% = The drug concentration (μ g/ml) that inhibited viral CPE by 50% calculated by using a regression analysis for semilog curve fitting.
- TC_{50} = (Cell) toxicity concentration 50% = The drug concentration (μ g/ml) that reduced cell viability by 50%.
- AI_{50} = Antiviral Index = A single point ratio of the antiviral and anticellular effect of the compound, calculated with 50% reduction values (calculated by dividing the TC_{50} by the IC_{50}).
- SI = Selectivity Index = A ratio calculated by dividing the TC_{25} by the IC_{50} (based upon 6 one-half-log₁₀ dilutions, μ g/ml, the maximum scale is 0-320).
- TAI = Total Antiviral Index = The area between the cytotoxicity and the antiviral curves (based upon a scale of 0-100%).

New Drugs with 25% Antiviral Reduction Levels: Of the 3895 actual single drug tests, 127 new compounds demonstrated marginal antiviral activity, having antiviral reduction values equal to or better than 25%. This represents around 3.3% of the test compounds being active at this marginal antiviral reduction level. In general, when compared to the 95% and 50% antiviral activity categories, the compounds in this (25%) category do not appear to have any significant antiviral promise and probably do not need to be confirmed any further.

4.1.1.5 Confirmatory Assays:

The CPE-inhibition assay procedure using (VR and/or MTT) was employed to confirm the inhibitory effects of compounds shown to be active in the primary screen. The results of the confirmatory evaluations are summarized on Table 5. Except for a few compounds (most of which showed only marginal to moderate activity), the antiviral effects of the compounds against VV were confirmed. Thirty-seven compounds have confirmed activity that is significantly greater than the positive control drugs as demonstrated by Selectivity Indices (SI) > 100 and high Virus Ratings or TAI's. Of these highly effective compounds, nine (AVS-1985, 2994, 3679, 3912, 4224, 4225, 4226, 4227, and 4533) have SI's of > 1000.

All of the 37 most active compounds should be considered for *in vivo* evaluation against VV. Furthermore, these compounds should be examined for activity against other DNA viruses such as Herpes Simplex virus, Varicella-Zoster virus, and Human Cytomegalovirus.

Table 5

Confirmatory Assays for Compounds Active Against Vaccinia Virus (VV)

<	u ⊢	•	•	+	+	+	+	+	+	٠	•	•	+	+	+	+	+	+	+	+	+	+	+	•	+	+	+	+	+	+	+	٠	+	+	
	Assay	4	TA	Ē	S S	3	CPE	363	8	CPE	300	E	345	343	HTT	HTT	SPE	HTT	HTTH	H	365	96	36	CPE	35	CPE	CPE	CPE	Se	MTT	H	CP.	E	H	
	I	10	72 52	8.7	2	1.90	1.70	1.10	5.	3, 10	7 80	65.56	5.40	3.50	17.58	11.42	1.20	26.53	21.87	18.09	4.20	3.30	1.40	0.60	4.70	4.30	5.20	5.50	6.30	29.69	45.74	4.00	18.85	23.11	
	15	2.10	11.81 >	5.66	3.8	2.60	3.10	8	3.30	11.40		_	100.00	4.10	4.36	1.91	1.20	5.87 >		3.77 >	4.60 >	3.90	1.80	9.	4.60	5.70	15.90	100.00	410.00	24.69 >	6.21 >	1.70 >	4.36 >	4.21	
	A1 %	8	200	88		9:	0.00	9	0.0	145.50	3 20 >	0.00	100.00	32.00	0.0	0.0	1.00	3.41	3.39	0.0	100.00	32.00	0.00	0.0	100.00	100.00	100.00	90.01	32.00	34.66	12.14	100.00	0.0	0.00	
	70 95	320.00	320 00	320.00	320.00 >	320.00 >	320.00	320 00	320.00	320.00	320.00 >	320.00	320.00 >	320.00	10.00	10.00	320.00 >	320.00 >	320.00 >	320.00	320.00 >	320.00	320.00	320.00	320.00 >	320.00 >	320.00 >	32.00	320.00	32.00 >	32.00 >	320.00 >	3.20	10.00	
	10 %	120.00	× 00 0		320.00 >	320.00 >	0.00	A 00 0	0.00	2.20	100.00	0.00	3.20 >	10.00 >	0.00	0.00	320.00 >	93.90 >	94.50 >	0.00	3.20 >	10.00	0.00	0.00	3.20 >	3.20 >	3.20 >	3.20	10.00		2.64 >	3.20 >	0.00	0.00	
	AI 50	200	11 81	ر ا	8.8 8.8	2.60	3.10	8	3.30	36.36	15.24	_	100.00	4.10		13.98	4.85	6.03	5.66	4.77	4.60		3.20	1.00	09.4	5.70	16.00	100.00	410.00	85.22	63.84	_	14.89		
	TC 50	320.00 >	320.00 >	320.00 >	320.00	320.00	100.00	100.00	320.00	320.00	320.00 >	320.00 >	100.00	3.20	10.00	10.00 >	320.00	320.00 >	320.00 >	320.00 >	3.20 >	3.20	320.00 >	320.00	3.20	3.20	10.00	32.00	100.00	32.00 >	32.00 >	32.00	3.20 >	9.56	
	10 50	100.001	27.10 >	47.40 >	81.00	28.00	32.00	112.00	98.00	8.80	10.00	3.11 >	1.00	0.80	0.55 >	0.71 >	00.99	53.10 >		67.10 >	0.70	0.80	100.001	320.00	0.70	0.60	0.60	0.30	0.20	0.38 >	0.50 >	0.32	0.21 >	0.19	
	AI 25	10.00	33.40	10.58	3.20	3.20	10.00	3.20	10.00	10.00	32.00	213.00	48.00	1.50	10.70	3.34	1.50	8.05	6.27	5.46	10.00	10.00	10.00	2.10	15.20	10.00	100.00	100.00	152.00	92.71	18.92	3.10	6.40	5.85	
	10 25	320.00 >	320 00	268.00		100.00	100.00		100.00	100.00	320.00 >	320.00 >	32.00	1.00	2.38	1.37	100.00	312.00	267.00	253.00	3.20	3.20	320.00	210.00	3.20	3.20	10.00	32.00	32.00	9.27 >	3.12	1.00	0.94	0.81	
	10 25	32,00 >	0.55 >	25.40	32.00 ~	32.00 -	10.00	- 10.00 -		~ 10.00	10.00 >	1.50 >	99.0	9.0	0.22	0.41	99.99	38.70	42.50	46.30	0.32	0.32	32.00	100.00	0.21	0.32	0.10		0.21	0.10	0.17		0.15	0.14	
	Oiff.	4	1.030	1.278	¥	ş	ş	4		V.	4	1.236	¥	¥	1.122	1.334	≨	1.122	1.247	1.439	¥	¥	4	ž	¥.	¥	¥	¥	¥	1.113	1.257	Y Y	1.122	1.367	
	<u>:</u> *	:	OVO	107	:	:	;	:	:	:	:	98K	:	:	OYX	50	:	0X2	305	Ĭ	:	:	i	i	i	:	:	;			110	:	o X	7	
	Test Date	07/31/86	06/14/90	07/19/90	07/24/86	05/56/88	07/17/86	02/04/88	05/26/88	06/12/86	07/17/86	08/17/89 08K	07/24/86	05/26/88	06/21/90	07/12/90	08/07/86	06/14/90	05/11/20	08/06/80	07/17/86	05/26/88	08/07/86	05/26/88	07/17/86	05/26/88	07/17/86	10/27/88	11/10/88	01/11/90	03/53/50	07/24/86	06/21/90 OYX	07/12/90	
	Ship-	8 8			20	416	10	24	416	7	4	5	05	416	29	29	8	29	67	29	5	416	95	9	5	416	5	99	95	82	29	20	29	29	
	AVS Ho.	** VIRUS W	1000	9	202	200	6200	200	6200	0138	0138	0138	0148	0148	0148	0148	9020	9020	9020	9020	0215	0215	1520	0257	2220	2220	0303	0303	0303	0303	0303	9790	9790	9646	

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	Assay	Type	ų.	L PE	, de			HTT	CPE		- L	SP.	CPE				2	343 CPE	HTT	H			CPE	MTT	HT	HT	MTT	HIT	HT	I	35	CPE	4		2	MIT	HTT	HTT	TT.	
							_		_		_					_		_				_		_	_										_					
		TAI	-	0.70	20		-	2.30	1.50			5.6	2.60	45.1	12 07		200	2.7	17.96	15.9		3.5	4.30	38.05	46.1	15.4	41.18	5.4	0.0	3.83	1.50	0.0	7.20		10.6	16.0	55.85	73.3	8	
		S	2.00	1.33	2.50		1.72 >	0.0	4.50	07 /	8	166.00	1.70	× 06.59	22 20 1	, 03.43	3	1.00	3.35	3.17 >		3.	32.00	12.10 >	13.80 >	1.98 >	13.50 >	1.08 >	00.00	0.00	4.10	0.00	313.00 >	6450 00	2650.00	2.65 >	. 72.50 >	> 177.92 >	. 5.59 >	
		AI 95	0	0.00	0.00		0.00	0.0	0.00	0	3	100.00	0.00	26.40		3 8	3.5	0.00	0.00	0.00	•		10.00	0.00		0.0	12.26	0.00	00.00	0.00	0.00	0.0	320.00 >1000.00	00 0000	320.00>10000.00	8.0	0.0	0.00	0.0	
		10 %	320.00	32.00	320.00		320.00	320.00	320.00	230 00	350.00	320.00	320.00	320.00 >	220 00			320.00	320.00	320.00	90	360.00	320.00	10.00	32.00 >	100.00	100.00			320.00	100.00	320.00	320.00	720 00	320.00	320.00	0.32	10.00	0.01	
		10 %	00 0	0.00	0.00		0.00	0.00	0.00	6	3.	3.20	0.00	12,10 >	42 00 ×	3.50	3.5	0.0	0.00	0.00	00	3.2	32.00 >	0.00	0.96	0.00	8.16 >	0.00	0.00	0.00	0.00	0.0	0.32 >	20 0	S 50.0	0.00	0.00	0.00	0.0	
		AI 50	8	1.33	2.50		1	0.0	4.50	40	8	8	1.70	8	2	3	8.9	1.00	2.00	8	8	3.	32.00	12.40	20	50	39.10	34	00	0.00	4.10	0.00	00	9	3 3	72	20	S	29	
		Y	•	-	~	i (N	o.	4		j	166.00		110	50 50	ξ .	•	-	'n			'n	3	. 12.	ĸ	9	ĕ		0	Ö	4	Ö	313.00	EVED	3650.00	7.24		> 312.50	ķ	
		TC 50	8	32.00	320.00		237.00	177.00	100.00	8	9.00	100.00	10.00	196,00	176.00	3 8	360.00	100.00	293.00	320.00 >		30.00	100.00	8	.17	.10	99.00	320.00 >	00	234.00	32.00	100.00	100.00	8	32.00	7.24	.35 >	10.00	6.	
		2	5	2	320		23/	14	100		3	5	2	5	176	2 6	2	9				2	5	2	0	47	8	320	247	ž	32	5	100	5	אַ י	_	0	유	0	
		10 50	16.90	24.10	129.00		85.50	0.00	22.00	50	22.00	0.60	5.70	3	87 2		3.5	100.00	58.60	< 07.89	8	3.	3.10	0.81 ×	0.36	67.7	1.69	238.00 >	0.00	0.00	7.80	0.00	0.32		L .	8.	0.00	0.03 >		
		52	8	2.10	3.30		4.59	1.32	6.60		2	8	8	8	2	2 9	3	6.60	5.23	8.4	,	9	8	8	20	6.11	30	19	00	0.00	9.60	9.60	1.00 <	5	3 :	2.65 <	8	> 26	07.6	
		¥	21.00	~	M		4		Ó	•	ń	313.00	21.00		02 57		Ö	•	Ŋ	4	•	Ó	8.0	145.00	20.	ø	30.30	-	0	0	ø	ø		7	5000	7	159.00	177.92	0	
		TC 25		21.00	~	,	147.00	82.00	99	77	8.8	100.00	9.60	100,001	2		V	99.00	196.00	217.00		9	86.00	9.74	5.02 >	8.87	22.80	257.00	173.00	167.00	21.00	99.99	0.32 >		21.00	2.65 >	0.35	2.69	0.01	
		10 25	1.20 -		66.00		32.00	62.20	10.00	2	9. L	0.32	0.32	00	1 71		25.00	10.00	37.60	43.50		3	1.00	0.07	0.10	1.45	0.73	154.00	00.00	0.00	3.20	10.00	0.32 <	200	0.00	9.	0.0	0.03	0.0	
		Diff.	5	4	4		1.31	1.206	NA AM		۱ ≦	NA A	¥.	1.384			Š.	×.	1.163	1.816		<	¥	1.7%	2.374 <	1.687	1.107	1.163	1.832	1.696	Y.	¥	¥			1.975 <	1.975	1.077 <	1.077	
	Ple	**	:	:	:		OSA	01L	:		:	:	:	Y	8			:	910	8		•	:	03E	040		OYE	910	OUR	3	;	:	;				5	0Y2	0Y2	
		Date	10/22/87	11/12/87	05/12/88		03/53/50	04/19/90	08/13/87	06/113/00	05/15/08	08/0/98	05/12/88	03/23/89	04/01/80	60/10/20	00/50/20	02/18/88	04/12/90	05/03/90 OUQ		10/12/00	10/27/88	05/04/89	06/01/89	05/31/90	06/21/90	04/12/90	05/03/90	05/54/90	08/27/87	05/12/88	07/24/86	08/76/30	99/97/60	02/23/89	02/23/89	06/21/90	06/51/90	
	Ship-	Ment	30	30	30		z	ઢ	27	22	,	27	22	25	2	; ;	2	33	8	t		ç	94	99	95	29	29	3	8	8	28	28	02	417	0	418	418	29	29	
	AVS		2070					1017	1159			35	1160	1214				1392	1644	1644							1654	1838	1838		1947	1947	1985						**1985	

* Based on combined results of plates 01M, Drug 1 and 01M Drug 2, the $1C_{50}$ is 0.00441 μ g/ml, the TC_{25} is 2.65 μ g/ml and the SI = 601.

** Based on combined results of plates 072, Drug 1 and 072, Drug 2, the $1C_{50}$ is 0.03179 $\mu g/ml$, the TC_{26} is 5.69 $\mu g/ml$ and the SI = 3179.

(U F	* *	+ +	* *	• •	+ + :	• •	+ +	* *	+ +	• •	• •	* *	• •	+ +	+ +
Assay Type	2 2	5 5	2 2	2 2	3 3	E	5 5	3 3 3 3 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5	2 2	F E	55	9 6 8 8	2 2	55	9 9
141	7.40	6.80 5.60	5.90	5.00	2.10	32.18	3.20	0.90	3.40	28.57	1.40	2.80	1.70	0.60	1.50
3	533.00 2	188.00	160.00	37.00	6.20	10.01	5.60	2.10	6.40	10.57	4.20	2.30	3.20	2.10	3.60
N 95	100.00	32.00 >	32.00 >	100.00	3.20	10.01	10.00	0.00	10.00	9.0	32.00	32.00	0.00	0.0	1.0
20 22	320.00 >	320.00 >	320.00 >	320.00 > 100.00 320.00 > 100.00	320.00	320.00	320.00	32.00	320.00 >	32.00	32.00	320.00	320.00	320.00 >	320.00 >
30 31	3.20 >	10.00 >	10.00 >	3.20 >	100.00	29.30	32.00	0.00	32.00 >	0.00	1.00	10.00	0.00	320.00 > 0.00 >	320.00 >
AI 50	533.00	188.00	160.00	7.70	6.20	18.04	5.60	3.20	6.40	78.06	2.00	2.30	3.20	7.10	3.60
10 50	320.00 >	320.00 >	320.00 >	10.00	100.00	242.00	32.00	10.00	32.00	32.00 >	3.20	10.00	320.00 >	320.00	320.00
10 50	0.60 \$	1.70 > 2.80 >	3.60 >	1.30	16.00	13.40	5.70	3.20	3.60	0.41 > 0.15	0.50	2.40	100.00 > 67.00 >	151.00	75.00
AI 25	1000.00	> 485.00 > 152.00	152.00	21.00	9.9	17.73	21.00	21.00	10.00 10.00	22.71 18.00	3.10	9.00	10.00	6.60	9.9
10 25	320.00 >1000.00 320.00 >1000.00	320.00	320.00 >1000.00 320.00 > 152.00	6.60	8.88	135.00	21.00	9.60	21.00	1.80	9.0	9.00	320.00 >	210.00	210.00
10 25	0.32 >	0.66 > 2.10 >	0.32 >	0.32	0.01	7.59	3.20	0.32	2.10	0.10	0.21	1.8	32.00 >	32.00	32.00
Diff.	· ·	\$ \$	11	\$ \$	MA MA	1.086	\$ \$	* *	\$ \$	0.967	* *	4 4	\$ \$	* *	\$ \$
ž=	11	: :	::	: :	: : 8	X	; ;	::	! - !	720 100	; ;	: :	::	: :	: :
Test P Date	07/24/86 -	07/24/86 -	07/24/86 -	10/16/86 06/16/88	12/04/86	06/21/90 0YK	11/13/86 -	11/13/86 -	12/04/86 -	06/27/90 02J 07/19/90 10H	12/18/86 -	12/11/86 05/12/88	12/11/86 - 05/26/88 -	12/11/86 - 05/12/86 -	02/12/87 05/12/88
Ship-	05 41b	416	416	98 41c	225	6 6	==	==	==	79	5 5	ដដ	13	a t	5 5
AVS No.	1986	1987	1988	2170	2275 2275	2275	2290	2291	2297 2297	2318 2318	2321	2323	2326	2336	5449

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Assay Type	FER	e e	9		TIM	S C	CPE TPE	HTT	CPE	CPE	CPE	GE	GE	CPE	CPE	CPE	CP.E	CPE	CPE	CPE	SP.	CPE	CPE	CPE	96 66	CPE	CPE	GP.	CPE	95 M	
1¥1	20.94	900	2.00	21.40	33.06	1.40	0.00	10.95	3.25	3.60	1.60	0.80	0.80	0.60	1.10	1.30	3.30	3.60	5.20	2.00	7.30	7.10	5.10	5.40	5.30	6.00	4.30	6.10	5.30	2.90	
2	3.79	6	18.00	50.7	6.86 >	3.30	1.80	2.07 >	3.03	5.65	7.70	0.50	2.00	09.0	4.50	3.20	29.00	32.00	136.80	89.20	2 00.000	> 786.00	133.60	00.00	139.03	206.50	50.20	207.00	134.10	81.90	
A1 95	0.00	9	1.00	10.65	3.38 >	0.00	0.00	0.00	1.00	32.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	1.00 >	10.00 >	3.20 >	\$20.00 >10	3.20 >	10.00 > 133.60	3.20 > 100.00	32.00 >	10.00 >	^	32.00 >	32.00 > 134.10	10.00 > 181.90	
25 25	32.00	420.00	320.00 >	320.00 >	100.00	32.00	32.00	81.10	320.00 >	320.00 >	100.00	100.00	100.00	100.00	320.00	320.00	320.00	320.00 >	320.00 >	320.00 >	320.00 > 320.00 >1000.00 >	320.00 >	320.00 >	320.00 >	320.00 >	320.00 >	320.00 >	320.00 >	320.00 >	320.00 >	
10 95	0.00	9			٨	0.00	0.00	0.00	320.00 >	۸	0.00		0.00	0.00		0.00	0.00	320.00 >	32.00 >	100.00	٨	100.00 >	32.00 >	100.00 >	10.00 >	32.00 >	٨	10.00 >		32.00 >	
A1 50	43.17	05.8			98.9	3.30	1.80	3.08		5.65	7.70	0.50	2.00	0.60	4.50	3.20		32.00	136.75	89.17		^ 786.00	133.60	100.00	139.00	206.48	50.20	207.00	134.10	181.90	
10 50	27.40 8.57	320.00	320.00	142.00	100.00	10.00	10.00	9.38	3.20	10.00	100.00	10.00	32.00	10.00	320.00	320.00	320.00 >	320.00 >	320.00 >	320.00 >		320.00 >	320.00 > 133.60	320.00 > 100.00	320.00 > 139.00	320.00 >	^	320.00 >	320.00 > 134.10	320.00 > 181.90	
10 50	0.63	28 00	18.00	17.00	14.60 >	3.10	5.60	3.05	1.08	1.71	13.00	20.00	16.00	18.00	71.00	100.00	10.70 >	10.00 >	2.34 >	3.59 >	0.32 >	0.41 >	2.40 >	3.20 >	2.30 >	1.55 >	6.37 >	1.55 >	2.40 >	1.76 >	
AI 25	13.39	10 00	21.00	5.56	10.45	9.60	3.10	3.93	3.18	9.60	20.60	99.0	2.10	99.0	3.18	3.18	48.50	48.50	152.40	152.40	0.00 >1000.00 <	523.80	152.40	320.00	152.40	485.00	48.50	485.00	152.00	320.00	
10.25	2.40	210 00	210.00	69.00	100.00	6.60	6.60	6.29	2.10	9.60	00.99	9.90	21.00	9.60	210.00	210.00	6.60 > 320.00 >	320.00 >	2.10 > 320.00 >	320.00 >	320.00 >1	320.00 >1523.80	2.10 > 320.00 > 152.40	320.00 >	2.10 > 320.00 > 152.40	320.00 >	320.00 >	320.00 >	2.10 > 320.00 > 152.00	320.00 >	
10 25	0.27	21.00	10.00	12.40	9.57 >	1.00	2.10	1.6	99.0	1.00	3.20 -	10.00	10.00	10.00	99.00	90.99	6.60 >	6.60 	2.10 >	2.10 > 32	0.32 > 320	0.21 >	2.10 >	1.00	2.10 >	0.66 >	6.60 > 320	0.66 >	2.10 >	1.00 ×	
Diff.	1.009	4	¥	1.044	696.0	¥	¥N	1.325	¥	¥	NA NA	¥	¥	Y.	¥	Y.	¥.	Y.	¥	¥	NA A	Y.	*	¥	Y.	Y.	¥.	Y.	¥	¥	
=	02K 101	:	:	OXI	170	:	:	986	:	1	1	:	:	:	;	:		;		:	-	:	:	:	1	:	:	:	:	:	
Test Date	06/27/90 02K 07/19/90 10I	05/14/87		06/20/90	06/22/90	06/25/87	05/26/88	08/17/89 08P	06/25/87	08/27/87	05/28/87	01/21/88		10/13/88		05/12/88		08/27/87	07/16/87	08/27/87		09/24/87		08/27/87	07/16/87	08/27/87		08/27/87	07/16/87	08/27/87	
Ship-	79		7			22	416	416	22				97			77	92			92	92		92			92	92		92		
AVS.	2503	2568	2568	2568	2568	2688	2688	2688	2700	2700	2716	2716	2716	2716	2786	2786	2873	2873	2874	2874	2875	2875	2879	2879	2888	2888	2889	2889	2890	2890	

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Assay	CPE	ų	ָ טַּ טַּ			CPE	CPE	SPE CPE	CPE	Je J	3	CPE	CPE	9	CPE	CP E	CPE	e E	340	G G	CPE	CPE	CPE	CPE	CPE	CPE	CPE	HT	CPE	<u>ي</u> و	
																											_				
TAI	5.90	1 70	10	27 20	27.30	5.80	6.60	6.00		3 3	4.85	4.40	3.40	7.00	4.50	2.20	3.20	6.20	5.80	5.90	5.80	6.10	6.30	2.30	2.30	2.20	1.60	25.66	1.60	0 9	18.73
2	86.90 159.50	2	8	2,00	07.01	251.30	317.72	295.20 >	192.80	100.00	367.80	10.79		100.00		8.70	33.00	310.00	180.80	220.00	180.80	313.70	311.11	4.90	4.90	4.40	2.50	7.25 >	4.10	9:	2.47
N 95	10.00 >	5	8 8	3	3	32.00 >	100.00	3.20 >	11.00 >	0.00	0.00	0.00	0.00	0.00	0.00	32.00	1.00	1.00	10.00 >	32.00 >	10.00 >	10.00	10.00 >	3.20	3.20	0.00	0.0	0.00	3.10	8.0	8.5
26 21	320.00 >	120 00	320.00 >	200.00		320.00 >	320.00 >	320.00 >	320.00 >	100.00	100.00	320.00	320.00	320.00	320.00	320.00	320.00 >	320.00 >	320.00 >	320.00 >	320.00 >	320.00 >	320.00 >	320.00 >	320.00 >	320.00	320.00	320.00	10.00	32.00	32.00
1c 95	32.00 > 10.00 >	2		. ,		^	A	100.001	A	00.00		0.00	^	0.00	^	100.00	320.00 >	320.00 >	32.00 > 3	10.00 >	32.00 >	32.00 >			100.001		^	· 00.0	3.20	0.00	· 8.0
								-										M													
AI 50	86.90 159.53	7	0	45.70	15.39	251.28	317.72	295.20		100.00	367.80	64.01	26.05	100.00		8.70	33.00	309.95	180.80	220.00	180.80	313.70		4.90	8.3	4.40	2.50	17.57	4.10	8:	7.7
TC 50	320.00 >	200 00	100.00	00 002	200.00	320.00 >	320.00 >	320.00 >	320.00 >	32.00	32.00	320.00 >	320.00 >	320.00	320.00 >	100.00	320.00	320.00 >	320.00 >	320.00 >	320.00 >	320.00 >	320.00 >	100.00	100.00	100.00	100.00	15.80	3.20	0.09	e.9
	A A					٨	A	٨	٨	ij		A						٨	^	٨	^	^	^								
10 50	3.70 >	87 00	115.00	10 60	. X	1.27	1.00 >	1.08 >	1.66 >	0.32	0.0	6.4	12.28 >	3.20	3.20 >	11.00	9.80	1.03	1.77	1.45 >	1.77 >	1.02	1.03	21.00	21.00	23.00	40.00	8.8	0.80	5.60	6.7
AI 25	152.00 320.00	9	1.00	25 74	10.33	485.00	0.66 > 320.00 > 485.00	0.32 > 320.00 >1000.00	320.00	00.99	318.20	320.00	100.00	210.00		10.00	31.80	485.00	320.00	485.00	320.00	0.66 > 320.00 > 485.00	485.00	9.60	6.60	10.00	3.10	17.65	09.9	2.10	6.13
×	88	5	2 2	2	3	6	8	, 00	9	9	8	8	^ 0	8	8	8	2	8	8	6	6	8	8	8	8	8	2	2	2	2 2	2
5	2.10 > 320.00 > 1.00 > 320.00 >	210 00	66.00			> 320.00 >	> 320.	> 320.(1.00 > 320.00 >	21.	21.00	1.00 > 320.00 >	> 320.(210.	1.60 > 320.00 >	99.99	210.00	> 320.(1.00 > 320.00 >	0.66 > 320.00 >	1.00 > 320.00 >	> 320.(0.66 > 320.00 >	99.99	98	00.99	99.00	65.	2.10	9.6	=
10 25	2.10	5	99	12 10	16.10	99.0	99.0	0.32	1.00	0.32	0.02	1.0	3.20	1.00	1.60	9.60	6.60	9.0	1.00	99.0	1.00	99.0	99.0	10.00	10.00	6.60	21.00	3.69	0.32	3.20	2
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Diff.	\$ \$	*	*	1 344	-	¥	¥	¥	¥	¥	¥	¥	¥	4	¥	¥	¥	¥	¥	4	×	¥	¥	¥	¥	*	¥	1.299	¥	¥ .	1.165
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Test	07/16/67	07/14/87	05/12/88	08/17/80 DDM	40/11/00	78/91/70	08/27/87	07/30/87	08/27/87	07/30/87	09/24/87	07/30/87	08/27/87	07/30/87	08/27/87	07/30/87	01/21/88	07/30/87	08/27/87	07/30/87	08/27/87	07/30/87	08/27/87	08/20/87	C5. 12/88	08/13/87	05/12/88	08/17/89 0BN	08/20/87	05/26/88	06/1/89 065
Ship- ment	22	*	2 2	2 %	07	97	56	92	92	56	92	56	92	92	97	92	92	56	92	92	97	92	92	27	27	27	27	27	22	419	9
AVS No.	2891	2804	2893	2802	5407	2895	2895	2896	586	2905	2902			2904		2906	5906	1162		2912	2912		2913	7967				9962			<u> </u>

< 0 ⊢	•	+	+	+	•	٠	•	+	+	•	+	+	+	•	+	+	•	+	+	+	+	+	+	+	+	+	+	+	+	*	•
Assay	CPE	CPE	CPE	CPE	CPE	CPE	CPE	CPE	CPE	CPE	CPE	CPE	CPE	CPE	CPE	CPE	H	CPE	CPE	CPE	CPE	CPE	CPE	CPE	S C P E						
14	2.10	1.30	7.10	8.60	8.70	5.20	4.50	2.50	2.90	7.10	7.10	6.80	7.00	2.30	2.50	1.60	47.60	3.20	3.40	4.20	4.10	4.00	4.20	8.00	7.20	7.20	9.80	9.10	8.50	1.40	1.10
8	9.50	3.20	916.80	3564.30	3564.30	3.20 > 104.50	> 47.00	5.50	18.00	> 653.60	320.00 > 102.70 >	100.00 > 433.70	× 671.40	5.80		5.59	× 17.68 ×	8.90	2.70	39.50	> 3.10	2.80	2.80	>1000.00	> 306.00	992.00	100001	3200.00	3200.00	2.90	2.00
A1 95	0.00	0.00	32.00	320.00 > 1000.00	320.00 >1000.00			3.20		10.00 >		100.00	100.00	1.00 >	1.00 >	10.00	. 10.36 >	0.00	3.20	0.00	100.00		100.00	320.00 > 320.00 > 1000.00	320.00 > 320.00	100.00	320.00 >10000.0	320.00 > 10000.0	>3200.00	0.00	0.0
75 %	320.00	320.00	320.00 >	320.00 >	320.00 >	320.00 >	320.00 >	320.00 >	320.00 >	320.00 >	320.00 >	320.00 >	320.00 >	320.00 >	320.00 >	320.00	320.00 >	320.00	320.00	320.00	320.00 >	320.00 >	320.00 >	320.00 >	320.00 >	320.00 >	320.00 >	320.00 >	320.00 >	320.00	320.00
10.88	0.00	0.00	10.00 >	0.32 >	0.32 >	100.00	100.00	100.00	320.00 >	32.00 >	1.00 >	3.20 >	3.20 >	320.00 >	320.00 >	~ 32.00 ~	30.90 >	0.00	100.00	0.00	3.20 >	3.20 >	3.20 >	1.00 >	1.00 \$	3.20 >	0.03 >	0.03 >	0.10 >	0.00	0.00
AI 50	9.50	3.20	916.80	3564.30	5815.00	104.45	46.81	5.50	18.00	320.00 > 653.60	320.00 >1028.70	320.00 > 433.70	671.40	5.80	6.50		17.68	8.90	2.70	M		2.80	2.80	1000.00	100.00 > 306.00	992.00	100.00 10000.00	3200.00	3200.00	2.90	2.00
TC 50	320.00	320.00	100.00	320.00	320.00	320.00 > 104.45	320.00 >	100.00	320.00	320.00	320.00 >	320.00 >	320.00 >	320.00 >	320.00 >	100.00	320.00 >	3.20	1.00	10.00	1.00	9.6	1.00	320.00 >1000.00	100.001	320.00	100.001	32.00	32.00	100.00	100.00
10 50	34.00	100.00	0.11	0.09	90.0	3.06 >	6.84 >	18.00	18.00	< 67.0	0.31 >	0.74 >	0.48 >	55.10 >	49.10 >	17.88	18.10 >	0.40	07.0	0.30	0.30	0.40	0.40	0.10	0.30 >	0.30	0.01	0.01	0.01	35.00	20.00
AI 25	10.00	3.20	2063.00	6563.00	6563.00	320.00 > 152.00	> 48.50	9.60	21.00	320.00 > 1000.00	320.00 >1000.00	0.32 > 320.00 >1000.00	320.00 >1000.00	100.00	10.00		23.78	656.00	2100.00	9.60	2.10 <	3.10	5.10	0.32 > 320.00 > 1000.00 <	>3200.00	2100.00	00.0099	6563.00	6563.00	2.10	2.10
TC 25	210.00	210.00	66.00	210.00	210.00	320.00	320.00	66.00	210.00			320.00		320.00	32.00 > 320.00 > 10.00		320.00 >	210.00	210.00	2.10	9.6	9.6	9.0	320.00	320.00	210.00	99.99	21.00	21.00	66.00	8.9
10.25	21.00	8.00	0.03	0.03	0.03	2.10 >	6.60 >	10.00	10.00	0.32 >	0.32 >	0.32 >	0.32 >	3.20 >	32.00 >	12.00	13.50 >	0.32	0.10	0.32	0.32	0.21	0.21	0.32 >	0.10	0.10	0.01	0.003	0.003	32.00	32.00
Diff.											v					1	1.299	~		٧	~			٧			٧				
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# #			2	2		2	2			2	2	2	2	2	2		90 6			2	2										:
Test Date	08/20/87	01/21/88	09/04/87	09/24/87	10/27/88	08/20/87	09/24/87	02/18/88	03/03/88	10/29/87	11/12/87	10/29/87	11/12/87	10/29/87	11/12/87	02/11/88	08/17/89 06%	03/03/88	03/17/88	10/22/87	11/12/87	01/21/88	05/26/88	01/14/88	02/04/88	02/18/88	01/21/88	02/04/88	02/18/88	01/14/88	02/04/88
Ship-	22	/2	27	27	3	27	27	33	33	31	31	31	31	31	31	32	32	32	32	32	32	32	416	32	32	32	32	32	32	32	32
AVS		24.62	7662		7662	5662		3513	3513		3533		3534		3535		3571	3586	3586				3593		3607	3607	3679		3679		3705
< 2	7	4	14	14	~	2	14	143	-	141	1-1	M	(La)	1	La.)	143	Lead.	I,J	144	~	M	M)	127	M	1	M	141	M	m	143	м

Table 5 (Cont'd)

(U F	+	+	•	•	•	+	+	+	+	+	+	+	•	+	+	•	•	+	+	+	٠	•	+	+	+	•	٠
Assay Type	5	CPE	CPE	CPE	CPE	CPE	CPE	TIM	CPE	CPE	CPE	CPE	CPE	CPE	CPE	SP.	CPE	CPE	CPE	365	340	5	CPE	CPE	GP.E	CPE	CPE
IAI	7.90	8.30	2.10	2.10	5.00	4.40	0.00	63.25	6.80	6.60	2.10	1.80	2.10	2.40	1.00	0.80	0.40	1.30	1.50	1.00	1.50	5.50	2.60	8.40	7.70	7.90	7.50
S	1075.00	1609.00	4.80	7.40	64.00	34.00	207.21		325.00	285.00		3.20	6.30	2.60	3.10	0.00	0.00	1.60	2.70	3.10	2.00	2.90	3.20	1760.00	996.00	000.00	707.00
A1 %	320.00 > 320.00 >1075.00	320.00 > 320.00 > 1609.00	0.00	1.00	10.00 >	10.00 >	31.25	0.00	100.00 > 325.00	100.00 >	1.00 >	0.00	3.20	10.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	000.000	>1000.00	320.00 > 100.00 >1760.00	320.00 > 100.00 > 996.00	320.00 > 320.00 >1000.00	320.00 > 100.00 > 707.00
70 95	320.00 >	320.00 >	320.00	320.00 >	320.00 >	320.00 >	100.00	320.00	320.00 >	320.00 >	320.00 >	320.00	320.00 >	320.00 >	32.00	32.00	10.00	320.00	320.00	320.00	320.00	320.00 >1000.00	320.00 >1	320.00 >	320.00 >	320.00 >	320.00 >
20 21	1.00 >	1.00 \$	0.00	320.00 >	32.00 >	32.00 >	3.20 >	0.00	3.20 >	3.20 >	320.00 >	0.00	100.00	32.00 >	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.32 >	0.32 >	3.20 >	3.20 >	1.00 >	3.20 >
A1 50	1075.00	1609.00	4.80	7.40	06.00		207.21 ~	211.00	325.00	285.00		3.20	6.30	5.60	3.10	0.00	0.00	1.60	2.70	3.10	2.00	2.90	3.20	1760.00	> 996.00	1000.00	707.00
TC 50	320.00 >1075.00	320.00 >1609.00	320.00 >	320.00	320.00 >	320.00 >	100.00	320.00 >	320.00 > 325.00	320.00 > 285.00	320.00 >	320.00 >	320.00	100.00	10.00	32.00	10.00	100.00	320.00	100.001	100.00	0.10	0.10	320.00 >1760.00	320.00 >	320.00 >1000.00	320.00 > 707.00
10 50	0.30 >	0.20 >	< 00'.29	43.00	5.00 >	9.50 >	0.48 >	1.52 >	1.00 >	1.10 >	88.00 >	100.00:	51.00	18.00	3.20	n. 00	0.00	64.00	118.00	32.00 >	> 00.09	0.03	0.03	0.20 >	0.30 >	0.30 >	0.50 >
AI 25	1524.00	3200.00	4.80	10.00	100.00	100.00		320.00	1000.00	1000.00	10.00	7.80	3.20	3.10	3.10	9.60	2.10	3.10	10.00	48.50	48.50	3.10	9.60	3200.00	1524.00	1524.00	1524.00
TC 25	320.00 >1524.00	320.00 >3200.00	320.00 >	210.00	320.00 >	320.00 >	100.00	320.00 >	320.00 >1000.00	320.00 >1000.00	320.00 >	320.00 >	210.00	00.99	9.60	21.00	6.60	90.99	210.00	320.00 >	320.00 >	0.07	0.02	320.00 >3200.00	320.00 >1524.00		320.00 >1524.00
10 25	0.21 >	0.10 >	66.00 ×	21.00	3.20 >	3.20 >	0.00		0.32 >	0.32 >	32.00 >	0.66 •	90.99	21.00	2.10	3.20	3.20	21.00	21.00	< 09.9	6.60 ×	0.05	0.01	0.10 >	0.21 >	0.21 >	0.21 >
Diff.	¥	4	¥	××	YN.	¥	¥	1.739 <	¥	KA	K	¥.	Y.	¥.	×	¥	¥.	¥.	¥	¥.	¥.	4	¥.	¥	¥	*	¥
ž *		;		:	:	:	;	01E	:	:	:	;		:	:	:	į	i	:	i	;	:	:	:	:		
Test	03/10/88	03/31/88	02/25/88	03/10/88	03/03/88	03/17/88	09/23/88	02/23/89	03/03/88	03/17/88	03/03/88	03/17/88	03/03/88	03/17/88	04/14/88		06/30/88	06/02/88	06/16/88	04/21/88	06/16/88	06/30/88	07/21/88	07/07/88	07/21/88	88/20/20	07/21/88
Ship-	×	×	×	×	35	35	35	35	35	32	35	35	35	32	37	37	37	41	17	39	39	11	77	75	75	75	75
AVS No.	3912			3913	3940			3940	3941	3941		3944		3963	4050		0507		4163	4182				4554	4554		4555

Table 5 (Cont'd)

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Assay Type	CPE	S S S	CPE CPE	GPE GPE	CPE	CPE	CPE	CPE CPE HTT	CPE CPE HTT	CPE CPE	9 6 6 6 6 6 6 6 6 6 6 6 6 6 6 6 6 6 6 6	CPE	CPE
141	8.30	8.70	1.50	1.20	1.50	5.30	5.90	1.40° 2.00° 10.89	1.70 1.40 12.70 2.75	9.20	6.00	6.10°	5.90
15	>1340.00 > 849.00	>3880.00	2.10	0.40	1.60	19.00	263.00 > 708.00	> 2.90 > 3.60 > 2.38 >	3.10	320.00 >3010.00 320.00 1930.00 0.00 0.54	1.00 * 180.00 * 180.00	> 206.00 > 174.00	32.00 > 160.00 32.00 > 180.00
A1 95	320.00 320.00	320.00 >1000.00 >3880.00 320.00 >1000.00 >4100.00	10.00	10.00	0.00	0.00	32.00	0.00	0.000		1.00 32.00 32.00	32.00 32.00	
70 95	320.00 >	320.00 >	100.00	100.00	320.00	320.00	320.00 >	320.00 320.00 320.00	320.00 320.00 320.00 320.00		320.00 × 320.00 × 320.00 ×	320.00 >	320.00 >
10 95	1.00 \$	0.32 >	10.00	10.00	0.00	0.00	10.00 >	0.00	0.00	1.00 ,	320.00 × 10.00 ×	10.00 >	10.00 \$
AI 50	1340.00	3880.00	2.10	0.40	1.60	19.00	263.00	2.90	3.10	3010.00 1930.00 0.54	180.00	206.00	160.00
1C 50	320.00 >1340.00 320.00 > 849.00	320.00	10.00	3.20	320.00 >	0.32	320.00	320.00 > 320.00 > 320.00 >	320.00 320.00 320.00 >	320.00 >3010.00 320.00 1930.00 32.00 0.54	320.00 × 320.00 ×	320.00 >	320.00 >
1C 50	0.20 >	0.08	5.30	8.10	202.00 > 176.00 >	0.02	1.20	110.00 × 89.00 135.00 ×	103.00 100.00 85.80 > 226.00 >	0.10 > 0.20	96.00 1.80 v	1.50 >	2.00 \$
A1 25	1524.00 1524.00	10000.00 3182.00	2.10	0.32	10.00	32.00	318.00	3.20	6.60	4848.00 2100.00 0.60	1.00	. 485.00	152.00
70 25	0.21 > 320.00 >1524.00 0.21 > 320.00 >1524.00	210.00 1	6.60	2.10	2.00 > 320.00 > 6.00 > 320.00 >	0.21	210.00 >	210.00 320.00 > 320.00 >	210.00 210.00 > 320.00 >	> 320.00 210.00 21.00	6.00 66.00 1.00 > 320.00 > 1.00 > 320.00 >	0.66 > 320.00 > 1.00 > 320.00 >	2.10 > 320.00 > 152.00 1.00 > 320.00 > 320.00
10 25	0.21	0.02	3.20	6.60	32.00 >	0.007	0.66	66.00 32.00 > 65.50 >	32.00 32.00 49.60 >	0.10	1.00	1.00	1.00
Diff.	\$ \$	\$ \$	11	\$ \$	\$ \$	\$ \$	5 5	NA NA 1.327	NA 1.735 2.243	11 1	\$ \$ \$	\$ \$	¥ ¥
# #	: :		::	: :	; ;	: :	; ;	: : 080	04M	::::	: ::	: :	; ;
Test	07/07/88 07/21/88	07/07/88	06/23/88 07/07/88	06/23/88 07/07/88	06/23/88	04/21/88	06/23/88	06/23/88 07/07/88 08/17/89 080	06/23/88 07/07/88 02/23/89 06/01/89		07/07/88 07/21/88 08/18/88	07/21/88 08/18/88	07/21/88 08/18/88
Ship- ment	45	75	75	75	75	39	45	222	7777	2 2 2 2	3 BB	43	43
AVS No.	4226	4227	6227	4230	4232	4275	4530	4531 4531 4531	4532 4532 4532 4532		4588 4619 4619	4620	4621

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Assay Type	CP.	FFF	9 5 5 5 9 9 9 9 9		EEE	EE	ĒĒ			EE EE	FFF
TAI	5.60	24.98 0.17 15.41	1.10	20.86	16.36 1.20 9.03	34.45	36.14	31.95	0.81	13.24 23.72 2.83 14.00	21.47 62.29 37.83
18	114.00	90.00	1.00 0.00 10.00 5.60	20.00 80.00	8.80 1.80 1.80	11.50	6.30 >	3.30	4.18 ×	6.48 > 0.22 > 0.48 > 0.48 > 3.47 > 3.47 > 0.48 > 0.	5.34 > 84.00 > 19.30 >
VI 95	32.00 >	0000	0.00	0000	0000	0.00	0.00	0.00	2.30	0.00 2	0.00 > 0.00 >
20 21	320.00 >	320.00 320.00 320.00	320.00 320.00 320.00 320.00 >	220.00 100.00 86.90	94.00 100.00 32.00	320.00	88.20	274.00 100.00 320.00 >	320.00	320.00 × 320.00 10.00	320.00 10.00 >
10 %	10.00 •	0000	0.00 > 0.00 > 320.00 >	0.00	0.00	0.00	6.00	30.10	139.00 \$	286.00 v 0.00 v 0.00 v	0.00
AI 50	114.00	28.40	1.00 0.00 10.00 5.60	0.00	3.90	21.70	12.90	9.00 7.82 5.34	1.90	3.06 6.48 3.47	12.20 133.00 42.70
10 50	320.00 >	277.00 297.00 275.00	320.00 > 320.00 > 320.00 320.00	50.50 23.70 23.90	53.80 23.30 20.60	320.00 >	24.00	66.40	182.00 320.00 >	320.00 > 320.00 > 10.00 >	12.20 > 10.00 > 13.70 >
10 50	2.80 > 1.80 >	6.00	320.00 > 0.00 > 32.00 > 57.00	11.20	13.80 0.00 8.06	14.70 >	1.61	8.50		105.00 * 49.40 * 1.00 2.88 *	1.00
AI 25	152.00	18.70 0.00 0.00	3.20 4.85 10.00	6.22 0.00 0.00	6.42	23.90	12.40	7.60		0.47 10.46 1.00 < 5.89	5.34 < 197.00 19.30 <
TC 25	320.00 >	96.80 188.00 187.00	320.00 > 320.00 > 210.00	18.80 14.70 15.40	39.50 15.20 14.30	169.00	10.20	39.60 32.80 320.00 >	68.60	22.90 320.00 > 1.00 ~	5.34 > 6.32 > 6.16 >
10 25	2.10 >	5.18 0.00 0.00	100.00 × 66.00 × 21.00	3.02	6.15 0.00 3.73	7.06	0.82	4.30	55.70 43.50	49.20 30.60 × 1.00 ×	1.00
biff.	\$ \$	1.753	1111	1.613 2.287 1.325	1.752 2.420 1.248	1.871	1.709	1.491	1.748	1.429 1.308 1.354 <	1.893 < 1.754 < 2.341 <
#		986	: : : :	10 % B	8 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	013 04A	2 × 0	90 00	035	08N 08M 076 08T	036 034 04P
Test	07/21/88 08/18/88	12/08/88 05/18/89 07/27/89	06/16/88 06/30/88 10/13/88 10/27/88	02/02/89 06/01/89 08/17/89	02/02/89 06/01/89 08/17/89	01/12/89 05/18/89	02/02/89 06/01/89	01/05/89 02/02/89 01/12/89		03/29/90 03/29/90 07/13/89 08/17/89	04/06/89 05/04/89 06/01/89
Ship- ment	53	8 8 9	3333	22.22	255	87	87	8 8 8 8	8 8 8	57 57 57	555
AVS S	7 7797	4822 4 4822 4 4822 4	7 7987	5 7£67 5 7£67 5 7£67	6636	5034 4	5035 4	5053 4		5138 5138 5146 5146 5146	5219 5 5219 5 5219 5
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Assay		ËË	ĒĒ	ĒĒ	ĒĒ	H	ĒĒ	ĒĒ	FE	ĒĒ	FF		ĒĒ	EE
Ţ	27.96 6.15 26.92	11.39	4.03	3.02	59.60 75.99	82.80	18.36	5.74	0.58	17.94 10.24	5.38	4.21 12.81 12.36 14.18	28.23	30.91
īs	8.21 × 0.00 6.44 ×	2.60 >	1.02 >	0.00	320.00 >	157.89 > 320.00 >	2.12 >	0.00	0.00	3.63	2.05 >	1.87 × 2.38 2.60 × 2.12 ×	8.39 > 3.15	3.11
A1 95	0.00	0.00	0.00	0.00	0.00 > 147.36 >	0.00 > 147.36 >	0.00	0.01	0.00	0.00	0.00		0.00	0.00
70 27	320.00 100.00 100.00	163.00	320.00	320.00	320.00	320.00	320.00	3.07	3.05	320.00 100.00	9.36	13.70 9.30 21.00 8.83	31.10	98.00
10 %	0.00	0.00	0.00	0.00	0.00 > 2.17 >	0.00 > 2.17 >	0.00	302.00	0.00	0.00	0.00	00000	0.00	0.00
AI 50	15.80 0.00 8.14	5.43	1.02	1.11	320.00	230.26	2.12	0.00	1.69	6.41	3.58	2.82 4.32 4.45	13.73	14.01
1C 50	98.10 31.40 100.00 >	43.80	320.00 >	320.00	320.00 >	230.00 > 320.00 >	320.00 >	1.87	1.69 >	37.90	3.58 >	2.82 × 2.71 3.15 2.29	20.10	27.30
10 50	6.20 0.00 12.30 ×	8.07 5.70	313.00 >	0.00 > 288.00 >	1.00	1.00	151.00 > 217.00 >	179.00	0.00	5.91	1.00	1.00 0.63 0.71 0.57	1.45	1.89
AI 25	3.78	4.38	1.8.1	1.21	320.00 < 320.00 <	157.89 < 320.00 <	3.41	1.21	1.00 <	5.23	2.05 < 2.26	1.87 × 4.32 × 5.08	12.20	13.21
10 25	50.90 15.20 79.10	20.60	320.00 >	320.00 >	320.00 >	158.00 > 320.00 >	320.00 >	1.21 >	< 1.00 - 0.85	21.50	2.05 >	1.87 > 1.49	12.20 >	17.50
10 25	2.17 4.02 3.39	3.82	177.00 > 256.00	264.00 >	1.00 •	1.00	93.90 >	< 1.00 0.23	4 1.00	4.10	1.00	< 1.00 0.39 0.42 0.24	1.00	1.32
Diff.	1.537 2.442 1.378	1.791	1.567	1.516	1.339 <	1.255	1.074 0.987	1.330	1.448	1.407	1.324	1.264 1.613 1.196 1.257	1.372	1.372
= *	9 038 9 04V 9 07C	9 032	90 08W	9 08x	9 070	9 042 9 07E	0 0xr	9 OKX	9 OKY 9 OMT	9 OLH 9 OMY	9 OF	9 0L5 9 0MV 0 0P8 0 0T1	9 OL8	9 0L8
Test	04/13/89 06/01/89 07/13/89	05/11/89 06/08/89	07/27/89	07/27/89 05/17/90	06/08/89 07/13/89	06/08/89 07/13/89	06/07/90	12/07/89	12/07/89 12/28/89	12/07/89	12/07/89	12/07/89 12/28/89 01/17/90 03/29/90	12/07/89 12/28/89	12/07/89
Ship- ment	222	22	28	28	28	28	19	2 5	2 2	33	19	2222	22	19
AVS	5450 5450 5450	5457	2488	2675	\$568 \$568	5569 5569	5601	5905 5905	5906 5906	5958 5958	5970 5970	5977 5977 5977 5977	5987 5987	5988 5988

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	Assay	3.5	HT	L !		L W	TH	T	F	HT	I	Ē	TTM	H	HTT	L	HT	E		H	TI	E	E:	Ē	TH	Ħ	Ē	TTM	TH.	T T	•	ËË	
	;	¥	2.90	16.42	20.22	26.11	21.79	16.66	15.43	7.66	\$1.01	5.70	24.55	13.24	33.52	20.53	19.67	14.60	3.	13.02	36.32	19.24	37.51	12.92	24.05	12.64	11.52	30.06	20.57	23.39		27.63	
	;	5	0.00	3.05		10.16 >	1.8		2.53	0.00		0.67 >	2.90			3.73	3.67	1.12 >		3.61	7.69 >	2.26 >	15.15 >	2.3	4.43 >	1.82 >		5.66 >		4.21		5.12 × 2.60 ×	
	;	VI 25	0.00	8.0	9.0	0.00	0.00	0.00	0.00	0.00	* 89	0.00	1.12 >	1.07 >	3.44 >	3.42	3.39	0.0	3	0.00	3.49 >	88	^	88	1.21 >		1.06 \$	3.39 >	3.41	8.8		8.8 8.0	
		7C 93	320.00	9.0	320.00	100.00	308.00	320.00	320.00	320.00	320.00 >	100.00	100.00	100.00 >	320.00 >	320.00 >	320.00 >	0.00	9.00	320.00	320.00 >	100.00	320.00	9.00	320.00 >	100.00	100.00 >	320.00 >		320.00		320.00	
		5 32 8 32	٨		٨	٨	0.0	٨	٨	0.00	^	0.00		^	٨		^	0.0	3		^	92.20 >				^	A	6.50 ×	٨	^ ^ 0.0	,	8.0	
		AI 50	0.00	2:5	45.10	159.38	3.51	7.06	5.60	0.00	10.05	1.07	2.99	1.90	49.9	5.56	5.58	1.23	3	5.51	69.7	2.2		3.46		1.62		2.66	5.21	6.22		5.12 3.90	
		10 SO	320.00	10.00 5.00 5.00		· 00.00L	172.00	234.00	233.00	313.00	320.00 >	100.00	100.001	100.00	320.00 >	286.00	316.00	100.00	3.3	320.00	320.00 >	100.00	320.00 >	100.00	320.00 >	100.00	100.00 >	320.00 >	275.00	283.00		320.00 >	
	:	10 50	0.00	1.76 >	3.5	0.63 >	49.10	57.70	41.60	0.00		93.50 >	33.40 >		48.20 >	3.5	8.8	\$1.00 ×		58.10	41.60 >	44.30 >	21.10 >	28.90 >		24.90 >	54.20 >	> 09.95	52.70	45.60		62.60 v 76.90	
	;	Q V	2.20	7.82	20.0	19.41	9.74	7.67	6.82	3.02	73.50	1.55	5.66	5.61	9.57	2.55	. a	9.06		5.68	13.94	5.5	25.46	5.24	99.9	5.46	2.51	7.52	68.7	13.01		7.15 4.03	
		S 21	2.20 >	5.35	71.0	0.3/	07.76	141.00	105.00	135.00	320.00 >	62.90	100.001	100.00	320.00 >	193.00	202.00	99.69	30.00	210.00	320.00 >	100.00	320:00 >	80.60	320.00 >	100.00	100.00	320.00 >	187.00	192.00		320.00 >	
		2 21		3.0	02.1	0.33	10.00	18.40	15.40	06.77	4 51.7	40.50	17.70 >	38.30 >	33.40 >	37.00	42.50	10.00	4.30	37.00	23.00 >	26.40 > 32.00 >	12.60 \$	15.40	^	٨	39.90 >	42.50 >	38.30	14.70		44.70 × 49.60	
		0111.	0.852 <	1.185	6 5	1.05/	0.921	1.189	0.660	1.160	1.039	0.938	876.0	1.277	1.079	1.025	. 165	1.021	-	1.183	1.209	1.265	1.210	1.313	1.137	1.313	1.798	1.143	1.1%	1.050		1.306	
	F.		6		3			8	ON O	OP N	005	SK K		010			5	88	3	8	8	55	8 8	OTE	700	OTE	90		OSK OSK	96	3	OPG OSL	
	Test		12/13/89	01/17/90	2/1/50	06/14/90	12/13/89	01/17/90	01/10/90	01/24/90	02/22/90	03/15/90	03/15/90	04/12/90	02/22/90	03/15/90	04/21/30	01/24/90	06/66/70	01/11/90	01/24/90	04/12/90	02/22/90	04/12/90	02/22/90	04/12/90	05/03/90	01/17/90	03/29/90	01/17/90		03/29/90	
	Ship-	Teor.	19	5 ;	8 :	8	61	5	62	62	3	3	62	8	3	3 :	8	25	7	29	3	33	33	2 2	62	3	62	62	3	33	\$	2 29	
	AVS		2665	2867	200	240	6042	6042	6145	6145	6193	6193	6500	6506	6210	6210	22	6213	2	6215	512	6217 6217	6218	6218	6219	6219	6219	6220	9550	6223	}	9259 9554	

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Assay Type	THE	EE	HTT	FF	TITE	HTH	H H	H III		FF		H H	FIN	T I	T I	E E	
TAI	19.07	6.02	32.30	23.24	79.64 0.00 0.00	22.77 30.67	18.78 12.91	3.48	8.58	7.85	15.73	9.46	21.69	3.50	6.74	11.28	
18	4.63 >	0.00	7.85 > 9.69	3.66	3.91	4.96 > 7.28	3.21	2. c. 2. g.	1.35	1.92 ×	2.40 > 1.05 >	1.67 > 0.00	3.89	1.24 > 0.00	1.16	1.8. 1.8.	
A1 95	0.00	0.0	33.56	0.0	0.00	0.00	0.0	0.00	0.0	0.00	0.00	0.00	3.45	0.00	0.00	1.06 v	
25 25	320.00	8.21	320.00 >	3.20	320.00 320.00 320.00	320.00	10.00	30.80	320.00 100.00	320.00 320.00	320.00 320.00	320.00 320.00	320.00 > 310.00	320.00	306.00	320.00 >	
56 22	0.00	0.00	9.54 >	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	92.70 × 0.00	0.00	0.00	302.00 >	
Af 50	0.00	2.85	18.66	8.03	5.81 0.00 0.00	17.92 22.51	6.45	2.71	3.15	2.67	2.40	1.67	5.65	1.49	2.18	78.1.	
10 50	320.00 >	2.13	61.10	1.86	251.00 303.00 320.00	78.20	8.93	19.50 17.70	89.20	228.00 233.00	320.00 > 320.00 >	320.00 > 272.00	264.00 162.00	320.00 > 233.00	152.00 143.00	320.00 >	
10 50	67.30 > 0.00	6.9 7.8	3.28	0.18	43.20 0.00 0.00 v	4.37	1.38	7.20	28.30	85.20	133.00 > 304.00 >	192.00 > 0.00	46.80 59.80	215.00 > 0.00	69.40 87.60	173.00 > 177.00 >	
AI 25	3.05	2.60	16.44	6.52	8.17 0.00 0.00	13.84	9.26	3.18	3.84	3.17	4.58	2.46	5.69	2.67	1.8 5.	2.50	
70 25	312.00	1.28	25.70	0.82	169.00 195.00 320.00	21.60	4.45	13.30	50.50	164.00	320.00 >	320.00 > 186.00	182.00 86.80	267.00	80.60	320.00 >	
10 25	36.20	0.49	1.56	0.13	20.70	1.56	0.48	4.16	13.20	51.70	69.90 >	130.00 >	32.00	100.00 88.50	40.50	128.00 >	
Diff.	1.154	1.096	1.341	1.193	1.248 1.215 1.278	1.194	1.305	1.146	1.039	1.275	0.956	0.872	1.202	1.440	1.436	1.299	
=	080	07.E	0SS OUZ	86	0SV 0T8 107	0XS 0ZH	107	105 105	101 101	111	108	970	109 117	020 11E	02X 11F	11F	
Test	03/01/90	04/12/90	03/29/90	04/19/90 0U4 05/10/90 0VV	03/29/90 (04/12/90 (07/19/90 1	06/07/90	07/19/90	06/21/90 0YL 07/12/90 105	06/21/90 0YT 07/12/90 101	08/02/90	06/27/90 02B 07/19/90 108	06/27/90 02G 07/19/90 108	07/19/90	07/12/90	07/12/90	07/12/90 02Y 08/02/90 11F	
Ship- ment	33	22	33	22	222	19	19	19	33	33	33	88	22	69	69	69	
AVS No.	6304	63%	6462	6467	6469	4219 6724	6754	1779	6783	6819	6973 6973	6983	9869	7003	7022	7023	

∢ ∪⊢	• •	+. •	* *	• •	• •	* * *	* *	* *	* * * * *	* *	* *	• •	• •	• •	+ +	
Assay	FF	FF	FF	FF	EE		FF	HI				H			HTT	
Ĭ	10.37	5.12	12.02	14.70 8.73	9.69	16.22 2.16 10.46	32.52	20.56	35.33 46.97 59.01 45.75 58.71	0.60	17.71	7.03	16.50	21.86	9.88	
15	2.3%	1.03	2.56	1.00 \$	0.00	3.00	7.19	3.18 .	8.87 × 27.28 × 29.68 × 21.30 × 41.98 ×	1.30 > 0.00 >	3.60 > 2.45 >	1.31 > 2.02 >	3.33	3.67	1.30 7.30	
AI 95	0.00	0.00	0.00	0.00	0.00	0000	0.00	0.00	124.04 > 48.88 114.81 > 13.19 11.65	0.00	0.00	0.00 >	0.00	0.00	0.00	
70 95	320.00 320.00	320.00	320.00	320.00	320.00	320.00 100.00 301.00	320.00 320.00	320.00	320.00 × 1 100.00 × 1 100.00 × 1 32.00 ×	320.00	320.00 320.00	320.00 >	320.00	320.00 100.00	320.00	
10.95	0.00	0.00	0.00	0.00	0.00	0000 0000 0000	0.00	0.00	2.58 v 0.87 v 2.73 v 2.75 v	0.00	0.00	315.00 >	0.00	0.00	0.00	
A1 50	3.36	3.15	4.45	0.00	2.92	67.43 0.00 3.29	17.28 37.25	66.6	95.47 137.85 312.50 100.00 48.43	1.30	3.60	1.31	3.04	5.53	15.78 6.78	
10 50	275.00	89.90	191.00	320.00 >	206.00	86.60 63.60 81.00	255.00 320.00 >	320.00 >	95.50 × 100.00 × 32.00 ×	320.00 > 320.00	320.00 > 320.00 >	320.00 >	190.00	308.00	254.00	
16 50	89.60	28.60	42.90	320.00 >	0.00	19.60	14.70	65.20 >	1.00 0.45 0.32 0.66 >	246.00 >	88.80 v	244.00 > 158.00 >	32.00	55.60	16.10	
AI 25	3.61	0.00	5.36 3.60	4.60	3.45	5.25	21.60	11.82	8.87 < 29.68 < 41.25	2.25	6.52	2.13	7.03	5.72	3.39	
10 25	187.00 177.00	29.40	110.00	320.00 > 286.00	136.00 179.00	58.80 28.30 47.00	106.00	207.00	8.87 × 12.30 × 9.50 × 6.82 27.70	320.00 > 249.00	320.00 > 245.00	320.00 >	107.00	204.00	3.20	
10 25	51.80	14.10	20.50	69.60 >	39.40	11.20 21.10 11.90	3.32	17.50	1.00 0.32 0.32 0.17	142.00 > 197.00	49.10 >	150.00 >	15.20	35.60	6.19	
biff.	1.185	1.513	1.080	1.018	1.277	1.253	1.186	1.372	1.372 * 1.004 * 1.115 * 1.238 0.973	1.221	1.239	1.296	1.230	1.354	1.241	
= =	16G 176	126	12A 167	151 168	142 174	358	116	111	15 X Z Z Z Z Z Z Z Z Z Z Z Z Z Z Z Z Z Z	177	178 193	52	17G 194	162	18N 10S	
Test P	10/11/90	08/08/80	10/06/90	10/18/90	08/30/90	09/20/90 11/01/90 11/29/90	10/04/90	10/04/90	08/02/90 08/23/90 10/04/90 11/01/90	11/01/90	11/01/90	11/01/90	11/01/90	12/13/90 1	01/10/91	
Ship-	22	22	22	22	22	222	£ £	E.E.	EEEEE	KK	23	23	KK	KK . C	22	
	70%	7 2017	7 0117	7302 7	7 1987	7413 7 7413 7 7413 7	7447	2 8772	7449 7 7449 7 7449 7 7449 7	7 8977	7 6977	7 6272	7 2827	8239 7		
No.	22	22	m n	KK	R R	222	22	77	22222	22	22	22	7.7	82	8355 8355	

Table 5 (Cont'd)

(∪ ⊢	• •	
Assay	THE	
1¥	0.00	
25	1.06 v	
A1 95	0.00	
70 95	320.00	
10.95	0.00	
A1 50	0.00	
10 50	320.00 > 192.00	
10 50	302.00 >	
A1 25	9.00	
70 25	80.20 > 320.00 > 0.00 88.70	
10 25	80.20 \$	
Diff.	1.091	
<u>:</u> *	9 4	
Test	01/10/91 180 01/31/91 10P	
Sh ip-	22	
AVS No.	8357	

This value is a virus rating (VR) rather than a TAL. The VR is a measurement of selective antiviral activity that takes into account the degree of inhibition of virus-induced CPE and the degree of cytotoxicity produced by the test compound similar to TAL is more accurate with MTT measurements.

The differential is the difference in the cell control and the virus control optical densities. DIFRNTL =

 (Viral) inhibitory concentration 25%, 50% and 95% = The drug concentration (kg/ml) that inhibited viral CPE by 25%, 50% or 95% calculated by using a regression analysis for semilog curve fitting. 1C25,50,96

= (Cell) toxicity concentration 25%, 50% and 95% = The drug concentration (µg/ml) that reduced cell viability by 25%, 50% or 95%. IC25,50,96 = Antiviral Index = A single point ratio of the antiviral and anticellular effect of the compound, calculated with 25%, 50% or 95% reduction values (calculated by dividing the TC_{25,50,95} by the IC_{25,50,95}). AI 25,50,95

Selectivity Index = A ratio calculated by dividing the IC₂₅ by the IC₅₀ (based upon 6 one-half-log₁₀ dilutions, #g/ml, the maximum scale is 0-320).

Total Antiviral Index = The area between the cytotoxicity and the antiviral curves (based upon a scale of 0-100%).

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Activity = A "+" denotes a rest that produced ≥25% reduction in CPE. A "-" denotes an inactive test (i.e. ≤25% reduction in CPE). ACT

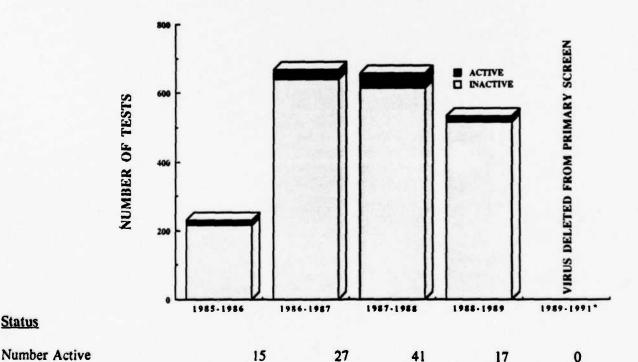
4.1.2 Adenovirus Type 2 (AD2)

The number of single drug tests carried out against AD2 during this contract period is summarized in yearly increments in Figure 11. During 1989, at the request of Ms. C. Susan Tiffany (Contract Specialist, Ft. Detrick, Frederick, Maryland), the Adenovirus was deleted for the primary screen.

A total of 2309 tests were performed during this contract period using a CPE-inhibition assay procedure. Adenovirus proved to be unsuitable for adaptation to the MTT assay without sacrificing sensitivity (Annual Report, December 15, 1988), therefore all screening with this virus was with the CPE-inhibition assay procedure. During the last year of AD2 testing (November 15, 1988 through November 15, 1989), the VR's of the positive control drugs ranged from 1.4-3.1 for Ribavirin and from 1.5 - 2.9 for Selenazofurin. The mean ID₅₀ of Ribavirin was 24.05 (± 11.1); the mean ID₅₀ of Selenazofurin was 12.35 (± 9.78). The selectivity of each of the positive control compounds, as measured by the TI, was: Ribavirin, mean 5.0 (± 4.1), median 3.9; Selenazofurin, mean 4.96 (± 3.21), median 3.91.

Out of the 2078 accepted single drug tests, 100 compounds demonstrated antiviral activity at greater than or equal to 50% reduction levels. This represents around 5% of the tested compounds having in vitro antiviral activity against the AD2 virus. The remainder, 1978 compounds (95%), were considered inactive with the CPE-inhibition assay protocol (Figure 11).

IN VITRO PRIMARY SCREEN: NUMBER OF COMPOUNDS FOUND ACTIVE AGAINST ADENOVIRUS DURING THE CONTRACT PERIOD



Five-Year

Totals

100

1978

2078

216

231

Status

Number Inactive

Yearly Total (Accepted

Single Drug Tests)

641

668

615

656

506

523

Represents 14-month period (November 15, 1989 - January 31, 1991) Figure 11

4.1.2.1 Confirmatory Assays:

The CPE-inhibition assay in HEp-2 cells was used to confirm the inhibitory effects of compounds shown to be active (i.e. a VR of ≥ 1.0 or $\geq 50\%$ reduction in CPE) in the primary screen. As shown in Table 6, 42 compounds demonstrated confirmed in vitro antiviral activity versus Adenovirus during this contract period. The activity of the remaining compounds (52/100) was either not confirmed on retest or there was insufficient drug available for retest. The compounds in Table 6 are presented in ascending numerical order. Data from the primary and the subsequent CPE-inhibition assays are listed chronologically for each compound. Some of the most effective compounds against AD2 were: AVS-2296, 2700, 2980, 2986, 3593, 4070, and 4167.

Table 6

Confirmatory Assays for Compounds Active
Against Adenovirus Type 2^a

AVS		Date					
No.	Shipment	Tested	<u>vr</u>	VR*	ID ₅₀	MTC	II
70	1	09/11/86	1.3	0.4	< 0.32	0.32	>1.0
	1	09/25/86	2.4	0.8	0.06	0.32	5.3
	1	09/15/89	0.65	0.22	-	1.0	••
79	1	09/11/86	1.3	0.4	2.3	10	4.4
	1	02/04/88	2.1	0.7	0.8	3.2	3.9
94	1	09/11/86	1.0	0.3	231.1	320	1.4
	1	04/21/88	1.0	0.3	222	>320	>1.4
136	4	09/25/26	1.3	0.4	96.9	320	3.3
	38	05/19/88	1.2	0.4	178.8	>320	>1.8
	41b	05/26/88	1.0	0.3	200	>320	>1.6
361	2	09/18/86	1.3	0.4	0.1	0.1	1.0
	2	10/30/86	0.6	0.2	0.3	0.1	0.3
	2	03/03/88	1.2	0.4	1.3	1.0	0.8
1215	27	08/13/87	1.1	0.4	62.9	320	5.1
	27	01/08/88	0.8	0.3	50.5	100	2.0
	27	01/21/88	1.3	0.4	53.7	320	6.0
	27	05/12/88	0.6	0.2	100	320	3.2
1337	33	02/04/88	1.5	0.5	88.4	320	3.6
	33	02/18/88	1.5	0.5	78.2	320	4.1
1984	2	09/15/85	0.35	0.12		320	••
	2	09/18/86	1.9	0.63	88	>320	>3.6
	2	05/26/88	0.7	0.23		> 320	**
2277	11	11/13/86	1.1	0.36	100	320	3.2
	11	05/12/88	0.35	0.12		320	
2296	11	12/04/86	1.2	0.4	10	32	3.2
	11	01/08/88	2.4	0.8	3.2	32	10.1
	11	01/21/88	2.0	0.7	10	100	10.0
2318	13	12/18/86	1.7	0.6	0.32	3.2	10.0
	13	09/04/87	1.8	0.8	3.59	10	2.79
	13	09/24/87	0.3	0.1	•	10	-
2350 ^b	14	11/05/87	1.8	0.6	6.3	20	3.2
	14	02/04/88	0.6	0.2	-	20	
	14	02/25/88	0.2	0.1	-	200	•
	14	05/26/88	0.6	0.2	63.1	200	3.2

Table 6 (Cont'd)

AVS		Date					
No.	Shipment	Tested	<u>VR</u>	VR*	ID_{50}	MTC	IL
2409	16	02/12/87	0.3	0.1		3.2	
	16	04/10/87	1.5	0.5	2.8	10	3.6
	16	01/08/88	1.1	0.4	2	10	5.0
	16	05/12/88	0.3	0.1		10	
	46	10/27/88	0.7	0.1	6.2	10	1.6
2563	21	01/14/00	1.1	0.4	46.0	100	2.1
2303		01/14/88		0.4	46.8	100	
	21 21	01/28/88 03/03/88	1.7 1.7	0.6 0.6	9.5 20.6	10 100	1.0 4.9
		03/03/00	•.,	0.0	20.0	100	4.5
2700	22	06/25/87	2.7	0.9	0.32	1.0	3.13
	22	07/09/87	3.7	1.23	0.21	3.2	15.20
	22	09/04/87	3.6	1.2	0.45	10.0	22.26
	22	09/24/87	1.8	0.6	0.45	1.0	2.23
2716	22	05/28/87	1.1	0.4	12.6	32	2.5
	22	01/21/88	1.4	0.5	14.1	100	7.1
	22	05/12/88	1.3	0.5	9.6	32	3.3
	46	10/13/88	0.9	0.3	10.0	32	3.2
2011	24	01/01/00	1.7	0.6	0.4	0.2	0.0
2811	24	01/21/88	1.7	0.6	0.4	0.3	0.9
	24	02/04/88	1.2	0.4	0.2	0.1	0.5
2812	24	02/25/88	1.1	0.4	0.1	0.1	0.7
	24	03/10/88	1.5	0.5	0.1	0.1	0.9
2960	27	08/13/87	1.2	0.4	44.4	100	2.3
	27	01/08/88	1.4	0.5	81.1	320	3.9
	27	05/12/88	0.8	0.3	•	320	-
2964	27	08/20/87	1.2	0.4	100.0	> 320	>3.2
	27	05/12/88	0.7	0.2	•	>320	•
2979	25	07/09/87	1.7	0.6	11.4	32	2.8
2717	25	01/08/88	0.9	0.3	68.4	320	4.7
	25			0.6	35.8	100	2.8
•	41b	01/21/88 05/26/88	1.8 1.3	0.6	10.0	32	3.2
2980	25	07/09/87	2.1	0.7	1.5	10	6.8
	25	10/29/87	2.0	0.7	1.0	3.2	3.3
	25	11/12/87	1.5	0.5	1.5	3.2	2.2
	25	05/12/88	1.2	0.4	5.0	10	2.0
2985	27	08/20/87	1.2	0.4	100	> 320	>3.2
	27	05/12/88		•	-	> 320	-
2986	27	08/20/87	1.7	0.6	100	320	3.2
2700	27	01/08/88	1.5	0.5	100	320	3.2
	21	01/00/88	1.3	0.3	100	320	3.2
2988	27	08/20/87	1.0	0.3	70.9	320	4.5
	27	01/08/88	0.6	0.2	-	320	•
	27	01/21/88	0.5	0.2	•	320	•
	27	05/12/88	0.9	0.3	307	>320	>1.0

Table 6 (Cont'd)

AVS		Date					
No.	Shipment	Tested	<u>VR</u>	VR*	ID ₅₀	MTC	II.
2994	27	08/20/87	1.5	0.5	25.5	25.5	3.9
	27	01/08/88	1.1	0.4	32.0	100	3.1
	27	01/21/88	0.7	0.2	64.1	100	1.6
3527	31	12/17/87	1.0	0.3	2.4	10	4.3
	31	05/12/88	0.4	0.1	•	100	•
3529	31	01/28/88	1.3	0.5	1.6	3.2	2.0
	31	02/18/88	1.6	0.5	1.8	10	5.5
3577	32	01/28/88	1.4	0.5	96.4	320	3.3
	32	02/18/88	1.8	0.6	113.0	320	2.8
3593	30	10/22/87	2.7	0.9	3.7	10	2.7
	30	10/27/87	2.7	0.9	3.7	10.0	2.7
	30	11/12/87	1.9	0.6	1.7	> 3.2	>1.8
	30	12/31/87	1.1	0.4	2.5	3.2.	1.3
	30	01/21/88	1.2	0.4	1.8	3.2	1.8
3610	32	12/31/87	0.6	0.2	100	320	3.2
	32	05/12/88	1.3	0.4	50.7	320	6.3
3705	32	01/14/88	1.7	0.6	73.1	320	4.4
	32	02/04/88	1.1	0.4	118	320	2.7
	32	02/18/88	0.9	0.3	147	320	2.2
	32	05/12/88	0.6	0.2	320	> 320	>1.0
3910	34	02/18/88	1.2	0.4	56.5	320	5.7
	34	03/10/88	1.1	0.4	204	>320	>1.6
4001	36	03/10/88	1.1	0.4	204	> 320	>1.6
	36	03/31/88	1.0	0.3	320	> 320	>1.0
4002	36	03/10/88	1.0	6.3	320	> 320	>1.0
	36	03/31/88	0.8	0.3		>320	
	36	04/14/88	0.3	0.1	-	>320	-
4004	36	03/10/88	1.2	0.4	193	> 320	>1.7
1001	36	03/31/88	0.8	0.3	173	>320	- 1.7
	36	04/14/88	0.5	0.2	-	> 320	
4041	36	03/17/88	1.1	0.4	320	> 320	>1.0
7071	36	04/21/88	1.1	0.4			
	30	04/21/86	1.1	0.4	320	>320	>1.0
4070	35	01/28/88	2.2	0.7	4.3	10	2.4
	35	02/18/88	2.3	0.8	3.8	32	8.5
4167	41	06/16/88	3.5	1.2	2.5	32	12.8
	41	06/30/88	2.3	0.8	6.6	32	4.8
4262	39	05/05/88	1.1	0.4	199	> 320	1.6
	39	05/26/88	0.6	0.2	-	>320	-
	39	06/16/88	0.5	0.2	••	> 320	-

Table 6 (Cont'd)

AVS		Date					
No.	Shipment	Tested	<u>VR</u>	VR*	ID_{50}	MTC	TI
4267	39	05/05/88	1.4	0.5	159	>320	2.0
	39	05/26/88	1.3	0.4	169	>320	1.9
4279	42	06/23/88	1.0	0.3	29.7	32	1.1
	42	07/07/88	1.0	0.3	40.3	100	2.5
4281	42	06/23/88	1.1	0.4	26.1	32	1.2
	42	07/07/88	1.1	0.4	37.6	100	2.7
4750	44	08/25/88	1.1	0.4	11.1	32	2.9
	44	09/01/88 .	0.4	0.1	8.4	10	1.2

a. Compounds are listed in ascending numerical order by AVS number. The results from the primary and the subsequent confirmatory CPE-inhibition assays are listed chronologically for each compound. Host cell were HEp-2. The VR is a measurement of selective antiviral activity which takes into account the degree of inhibition of virus-induced CPE and the degree of cytotoxicity produced by the test compound, determined by a modification of the method of Ehrlich et al. (Ann. N.Y. Acad. Sci. 130:5, 1965). VR* is the designation for the virus rating calculated by the method of Sidwell and Huffman (Appl. Microbiol. 22:797, 1971). The drug concentration which reduced the CPE by 50% (50% inhibitory dose, ID_{50}) was calculated using a regression analysis program for semilog curve fitting and is expressed as $\mu g/ml$. The minimum cytotoxic drug concentration (MTC) is also expressed as $\mu g/ml$. The TI of a test compound was determined by dividing the MTC by the ID_{50} .

b. The ID₅₀ and MTC for AVS-2350 are expressed as units/ml.

4.1.3 Yellow Fever Virus (YF):

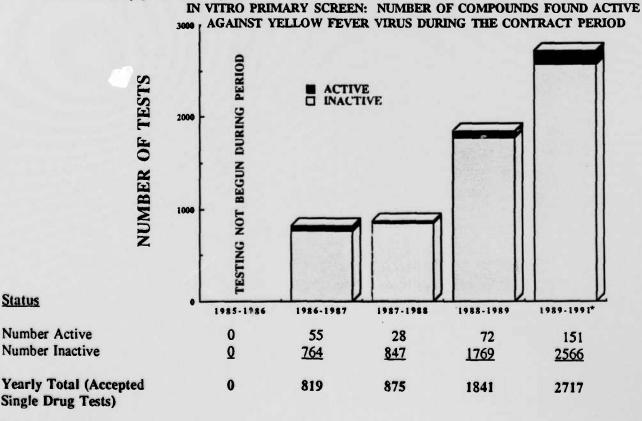
The number of single drug tests carried out against YF during this contract period is summarized in yearly increments in Figure 12. During this five-year period two main in vitro antiviral assay protocols were implemented:

- A standard CPE inhibition assay by virus rating (VR) (Annual Report, December 15, 1. 1988, Section 3.2.4).
- 2. Since November, 1988, MTT based-antiviral assay format.

A total of 7654 tests were performed during this contract period using both assay types. Routine testing was changed to the MTT-assay format to improve the efficiency and quality of the primary screening program in addition to being more cost-effective. Selenazofurin (AVS-0253) was tested in each standard virus rating (VR) CPE-inhibition assay as a positive control compound. Results of the positive controls (VR tests) were used as a guideline to assess the quality of each assay.

After the testing was converted to the MTT-assay format, we performed a total of 211 control compound assays with Selenazofurin during the last 26 months of the contract period. During this time 585 tests were internal (+++) virus load, cell load, and other quality control tests. Three hundred fifteen (315) tests were considered unsatisfactory based on the criteria of the quality controls set during this reporting period. The rest, totaling 2717 were actual single drug MTT-assays. The total number of MTT-assays (5960) tested during the last two years represents a 252% increase (improvement) in the total testing output as compared to the total of 1694 tests performed during the first 3 years of this contract.

Out of the 6252 accepted single drug tests, 306 compounds demonstrated antiviral activity at greater than 50% reduction levels. This represents around 5.0% of the tested compounds having in vitro antiviral activity against YF-virus. The remainder, 5946 compounds (95%), were considered inactive with both assay protocols (Figure 12).



Five-Year

Totals

306

5946

6252

Status

Represents 14-month period (November 15, 1989 - January 31, 1991) Figure 12

- 4.1.3.1 <u>YF-Quality Controls:</u> Two positive control compounds (Selenazofurin and 2-Thio-6 Azauridine) were used in the daily MTT assay sets as antiviral activity quality controls. The antiviral performance of the unknown compounds is compared to that of the positive control compounds. Compounds with equal to of better antiviral potency are considered active and are worthy of further *in vitro* profile studies and *in vivo* testing.
- 4.1.3.1.1 <u>Antiviral Activity of Selenazofurin vs YF Virus:</u> A summary of the antiviral and cytotoxicity performance of the primary control compound, AVS-0253 (Selenazofurin) is presented in Figure 13-A for 211 tests performed during November, 1989 through January, 1991.

Control Compound Antiviral Performance: Selenazofurin (AVS-0253) has been the sole control compound against YF in these MTT-assay screens. The mean and median antiviral inhibition and cytotoxicity patterns of the positive control drug (Selenazofurin) are illustrated in Figure 13-A.

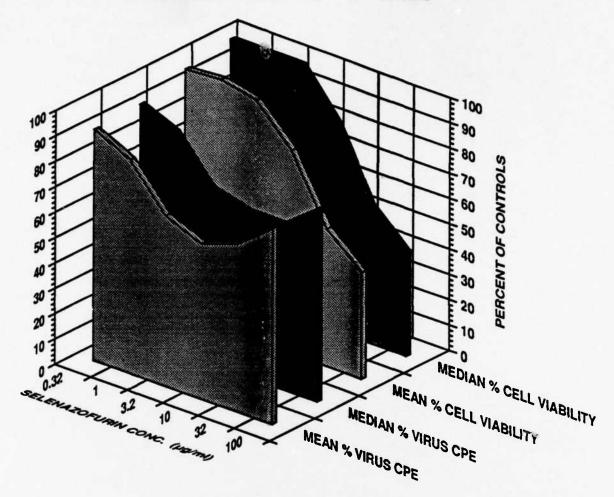
The 211 control tests performed with Selenazofurin gave a mean Total Antiviral Index (TAI) of 11.56% (SD \pm 9.12) and the median value was 9.32%. The TAI measures the overall antiviral effectiveness of the compound and it ranged from ~ 0 - 48% during this period. The mean Selectivity Index (SI) was only 1.06 (SD \pm 2.72) and the median SI value was 0, indicating poor antiviral selectivity for Selenazofurin and it ranged from ~ 0 - 19 during this period. However, the closeness of the mean and median values indicate that the present execution of the SOP is consistent and repeatable.

The mean Antiviral Index 25% (AI₂₅) value was 8.61 (SD \pm 15.13). The median AI₂₅ value was 3.76 (range 0.30 - > 125). The mean Antiviral Index 50% (AI₅₀) was 4.49 (SD \pm 11.54) with a median of 0 (range 0 - 70.86). This indicates that Selenazofurin does not consistently reach 50% antiviral reduction levels. The Antiviral Index 95% (AI₉₅) was not attainable with Selenazofurin versus Yellow Fever Virus.

The mean Antiviral Inhibitory Concentration 25% (IC₂₅) was $5.62~\mu g/ml$ (SD $\pm~5.84$). The median IC₂₅ value was $4.25~\mu g/ml$ (range = $<0.32~-32.00~\mu g/ml$). The mean Antiviral Inhibitory Concentration 50% (IC₅₀) was $2.49~\mu g/ml$ (SD $\pm~7.57$). The median IC₅₀ value was $0~\mu g/ml$ (range $0~-83.6~\mu g/ml$). This discrepancy indicates that the control compound Selenazofurin does not consistently reach 50% reduction levels. The mean Antiviral Inhibitory Concentration 95% (IC₉₅) could not be attained with Selenazofurin versus Yellow Fever Virus.

The average maximum antiviral inhibitory level of 211 Selenazofurin tests (Figure 13-A) was reached at 10 μ g/ml of the compound with 20% antiviral effect. Further increase of the drug concentration does not improve its antiviral activity. Maximum antiviral effect (~40%) was found with a simultaneous ~20% cytotoxic suppression. Above 10 μ g/ml concentration of the antiviral protection levels off to 20% of reduction level at 100 μ g/ml while simultaneously Selenazofurin becomes maximally toxic (~55%)

SELENAZOFURIN - VS -- YF VIRUS



CONCENTRATION (µg/ml)

% Viral CPE	% Cell Viability
,, ,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,	

Conc.(µg/ml)	0.32	1	3.2	10	32	100	0.32	1	3.2	10	32	100
Mean	91	84	67	61	66	77	97	98	95	80	54	42
Median	92	86	69	65	65	76	100	100	100	82	54	41
Std. Dev.	0.07	0.10	0.15	0.14	0.11	0.12	0.05	0.04	0.07	0.14	0.11	0.09

Figure 13-A

Average Antiviral and Cytotoxicity Values for 211 Positive Control Compound Tests

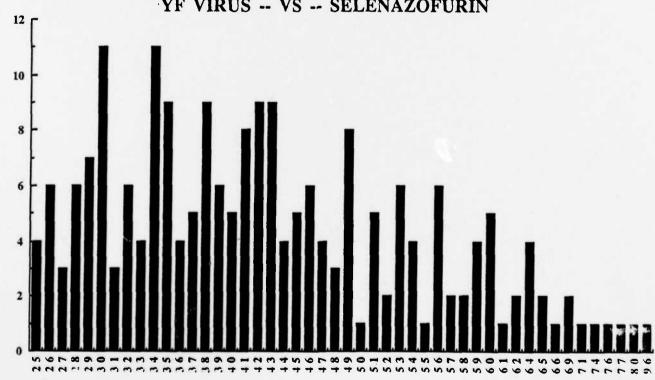
4.1.3.1.2 <u>Maximum Antiviral Effect of Selenazofurin vs YF Virus:</u> Since the metabolic activity of the cells was an unknown function during the testing period, it was monitored indirectly by measuring the maximum antiviral effect of the control compound Selenazofurin. This demonstrated the amount of infectious virus that was produced by the cells (Maximum Percent CPE).

A bar graph scatter plot (Figure 14-A) depicts the distribution of the maximum antiviral reduction values of all 211 control compound assays for Selenazofurin. The results indicate that the average maximum antiviral reduction obtained with the present SOP is around 43% (SD \pm 12.46) reduction levels. The maximum reduction levels vary from 25 - 96% but remain quite consistently around the median of 41%. The assay control values give a relatively broad bell-shaped distribution curve. This indicates quite a consistent day-to-day performance of the control compound in the YF-MTT assay.

During this period the positive control compound performance criteria for Selenazofurin versus the YF Virus was set at 25% reduction level. All assays in which Selenazofurin did not meet this accepted quality control level (>25%) were rejected (i.e., 315 unsatisfactory tests).

Since Selenazofurin is only marginally active against YF virus, better quality control compounds are needed. However, regardless of the poor performance of the YF quality control drug Selenazofurin, around 430 different compounds have equal or better antiviral activity against YF virus than AVS-0253. Some of these could certainly be used as a better *in vitro* antiviral control compound in this large-scale antiviral screening program.





FREQUENCY

PERCENT CPE REDUCTION

Figure 14-A

Maximum Antiviral CPE Reduction (%).

Summary of 211 Control Tests.

4.1.3.1.3 Cellular Cytotoxicity of Selenazofurin vs YF Virus:

YF-Control Compound-Cytotoxicity Performance: The 211 cytotoxicity values of the positive control compound Selenazofurin are also very consistent. The mean cell Toxic Concentration 25% (TC₂₅) was 16.49 μ g/ml (SD \pm 10.45) and the median was 15.80 μ g/ml (range of 0.24 - 62.9 μ g/ml). The mean cell Toxic Concentration 50% (TC₅₀) value was 57.38 μ g/ml (SD \pm 29.85) and the median was 54.70 μ g/ml (range of 8.51 - 100 μ g/ml). The mean cell Toxic Concentration 95% (TC₉₅) value cannot be attained with Selenazofurin versus Yellow Fever Virus.

As can be seen from Figure 13-A, the toxicity starts to become measurable above the concentration of 3.2 μ g/ml and the maximum toxicity has not been reached at 100 μ g/ml.

When the cytotoxicity reaches around 20% (10 μ g/ml), the control compound (Selenazofurin) loses its maximum antiviral effect (~35%). Above 32 μ g/ml the antiviral protection of Selenazofurin starts to decrease (down to ~20%). Selenazofurin becomes maximally toxic at 100 μ g/ml concentration. The highest selenazofurin concentration tested in these assays was 100 μ g/ml.

Selenazofurin has a definite cytotoxic suppression on cellular metabolism and growth. The TC_{25} and TC_{50} toxicity can be achieved with relative consistency at 100 μ g/ml.

4.1.3.1.4 YF-Assay Plate Quality Controls: Cell Load and Virus Load Parameters (Selenazofurin): The MTT assay is fundamentally dependent upon the quality of the assay plates. Our large-scale antiviral testing is dependent upon the uniformity of the test plates produced for the daily assays. Equal numbers of cell load and virus load as well as the consistent performance of the reagents used daily was monitored. A sample of the plate variation control for the period of November, 1989 through January, 1991, is presented in Figures 15-A, 16-A and 17-A.

YF-Control Compound-Cell Load Performance: A bar graph scatter plot of the mean cell control (O.D. reading) of 211 control assays is plotted in Figure 15-A. The results indicate that the cell O.D. readings reached a mean 1.130 (SD \pm 0.180) with a median of 1.140 (range of 0.400 - 1.680). This indicates that a uniform and equal number (18,000 cells/well) of cells are being loaded into every well in the 96-well plate during the day-to-day operation. The cells reduced MTT to formazan giving maximum blue color uniformly and consistently.

YF-Control Compound-Virus Load Performance: A bar graph scatter plot of the mean virus load O.D. readings of the 211 control assays is presented in Figure 16-A. The results indicate that the average virus load O.D. reading is 0.230 (SD \pm 0.110) with a median of 0.210 (range of 0.01 - 0.530). This demonstrates that a good cell destruction is taking place and a uniform load of virus (32 TCID₅₀) is administered on the cell monolayer with very consistent viral CPE results.

YF-Control Compound-Assay Differential Performance: A bar graph scatter plot of the mean O.D. differential values of the 211 control assays is provided in Figure 17-A. The results indicate that the average differential O.D. reading is 0.900 (SD ± 0.173) with a median of 0.891 (range 0.389 -1,440). The single bell-shaped curve is reasonably sharp and uniform. This reflects that the assays are executed consistently and are repeatable during day-to-day operation with close to 90% measurement accuracy.

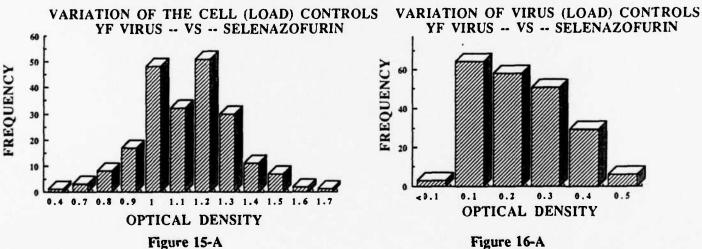
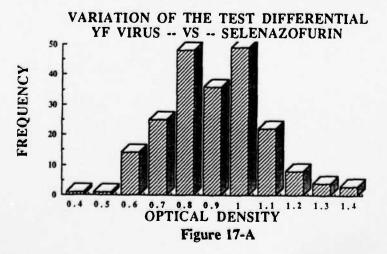


Figure 16-A



4.1.3.1 YF-Ouality Controls:

4.1.3.1.1 Antiviral Activity of 2-Thio-6-Azauridine vs YF Virus: A summary of the antiviral and cytotoxicity performance of the second control compound, AVS-6724 (2-Thio-6-Azauridine) is presented in Figure 13-B for 42 tests performed during November, 1989 through January, 1991.

Second Control Compound-Antiviral Performance: 2-Thio-6-Azauridine (AVS-6724) has been tested as a possible second control compound against YF in these MTT-assay screens. The mean and median antiviral inhibition and cytotoxicity patterns of this second positive control drug are illustrated in Figure 13-B.

The 42 control tests performed with 2-Thio-6-Azauridine gave a mean Total Antiviral Index (TAI) of 27.90% (SD \pm 14.50) and the median value was 25.00%. The TAI measures the overall antiviral effectiveness of the compound and it ranged from \sim 6.25 - 58.77% during this period. The mean Selectivity Index (SI) was 8.60 (SD \pm 8.90) and the median SI value was 5.20, indicating moderate antiviral selectivity for 2-Thio-6-Azauridine and it ranged from \sim 0 - 37.08 during this period. However, the closeness of the mean and median values indicate that the present execution of the SOP is consistent and repeatable.

The mean Antiviral Index 25% (AI₂₅) value was 21.50 (SD \pm 23.70). The median AI₂₅ value was 11.00 (range 2.18 - 98.96). The mean Antiviral Index 50% (AI₅₀) was 29.20 (SD \pm 30.30) with a median of 23.20 (range 0 - 162.60). This indicates that 2-Thio-6-Azauridine does not consistently reach 50% antiviral reduction levels. The Antiviral Index 95% (AI₂₅) was not attainable with 2-Thio-6-Azauridine versus Yellow Fever Virus.

The mean Antiviral Inhibitory Concentration 25% (IC₂₅) was 2.90 μ g/ml (SD \pm 3.40). The median IC₂₅ value was 1.50 μ g/ml (range = 0.32 - 17.50 μ g/ml). The mean Antiviral Inhibitory Concentration 50% (IC₅₀) was 6.50 μ g/ml (SD \pm 9.00). The median IC₅₀ value was 3.30 μ g/ml (range 0 - 49.30 μ g/ml). This discrepancy indicates that the control compound 2-Thio-6-Azauridine does not consistently reach 50% reduction levels. The mean Antiviral Inhibitory Concentration 95% (IC₉₅) could not be attained with 2-Thio-6-Azauridine versus Yellow Fever Virus.

The average maximum antiviral inhibitory level of 42 2-Thio-6-Azauridine tests (Figure 13-B) was reached at 10 μ g/ml of the compound with 70% antiviral effect. Further increase of the drug concentration does not improve its antiviral activity. Maximum antiviral effect (~77%) was found with a simultaneous ~15% cytotoxic suppression. Above 10 μ g/ml concentration the antiviral protection levels off to 50% reduction level at 320 μ g/ml while simultaneously 2-Thio-6-Azauridine become maximally toxic (~70%).

2-THIO-6-AZAURIDINE -VS- YF VIRUS

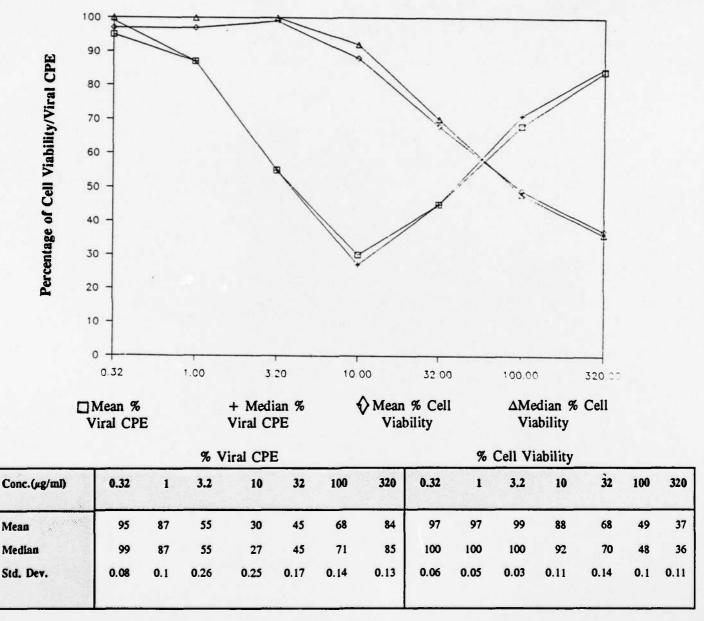


Figure 13-B
Average Antiviral and Cytotoxicity Values for 42 Positive Control Compound Tests

4.1.3.1.2 <u>Maximum Antiviral Effect of 2-Thio-6-Azauridine vs YF Virus:</u> Since the metabolic activity of the cells was an unknown function during the testing period, it was monitored indirectly by measuring the maximum antiviral effect of the control compound 2-Thio-6-Azauridine. This demonstrated the amount of infectious virus produced by the cells (Maximum Percent CPE).

A bar graph scatter plot (Figure 14-B) depicts the distribution of the maximum antiviral reduction values of all 42 control compound assays for 2-Thio-6-Azauridine. The results indicate that the average maximum antiviral reduction obtained with the present SOP is around 77% (SD \pm 17.20) reduction levels. The maximum reduction levels vary from 43 - 100% but remain quite consistently around the median of 76%. The assay control values give a relatively shifted bell-shaped distribution curve. This indicates quite a consistent day-to-day performance of the control compound in the YF-MTT assay.

Recommendations:

Based upon the data obtained in parallel studies with Selenazofurin, we recommend that 2-Thio-6-Azauridine (AVS #6724) will be used as a second control compound against YF virus. It's overall performance is much better than the present control, Selenazofurin. It is readily available from Sigma Chemical Company, it is inexpensive and works as effectively at low drug concentrations as Selenazofurin.

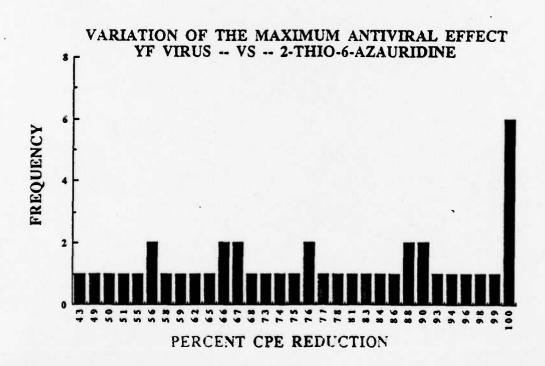


Figure 14-B

Maximum Antiviral CPE Reduction (%).

Summary of 42 Control Tests.

4.1.3.1.3 Cellular Cytotoxicity of 2-Thio-6-Azauridine vs YF Virus:

YF-Control Compound-Cytotoxicity Performance: The 42 cytotoxicity values of the positive control compound 2-Thio-6-Azauridine are also very consistent. The mean cell Toxic Concentration 25% (TC₂₅) was 29.80 μ g/ml (SD \pm 21.90) and the median was 26.80 μ g/ml (range of 7.73 - 100 μ g/ml). The mean cell Toxic Concentration 50% (TC₅₀) value was 95.80 μ g/ml (SD \pm 61.00) and the median was 94.30 μ g/ml (range of 26.00 - 320 μ g/ml). The mean cell Toxic Concentration 95% (TC₉₅) value cannot consistently be attained with 2-Thio-6-Azauridine versus Yellow Fever Virus.

As can be seen from Figure 13-B, the toxicity starts to become measurable above the concentration of 10 μ g/ml and the maximum toxicity has not been reached at 320 μ g/ml.

When the cytotoxicity reaches around 10% (10 μ g/ml), the control compound (2-Thio-6-Azauridine) loses its maximum antiviral effect (~77%). Above 10 μ g/ml the antiviral protection of 2-Thio-6-Azauridine starts to decrease (down to ~15%). 2-Thio-6-Azauridine becomes maximally toxic at 320 μ g/ml concentration. The highest 2-Thio-6-Azauridine concentration tested in these assays was 320 μ g/ml.

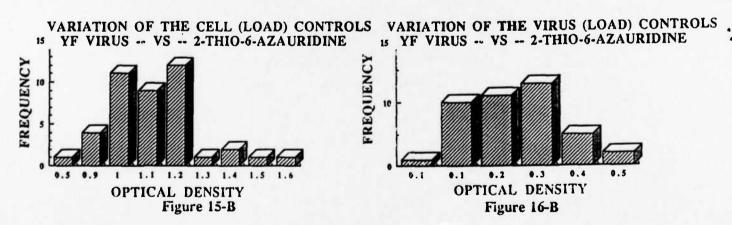
2-Thio-6-Azauridine has a definite cytotoxic suppression on cellular metabolism and growth. The TC_{25} and TC_{50} toxicity can be achieved with relative consistency at 100 μ g/ml.

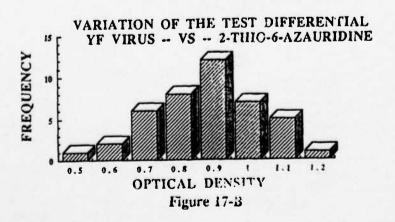
4.1.3.1.4 YF-Assay Plate Quality Controls: Cell Load and Virus Load Parameters (2-Thio-6-Azauridine): The MTT assay is fundamentally dependent upon the quality of the assay plates. Our large-scale antiviral testing is dependent upon the uniformity of the test plates produced for the daily assays. Equal loads of cell load and virus load as well as the consistent performance of the reagents used daily was monitored. A sample of the plate variation control for the period of November, 1989 through January, 1991 is presented in Figures 15-B, 16-B, and 17-B.

YF-Control Compound-Cell Load Performance: A bar graph scatter plot of the mean cell control (O.D. reading) of 42 control assays is plotted in Figure 15-B. The results indicate that the cell O.D. readings reached a mean 1.110 (SD \pm 0.180) with a median of 1.100 (range of 0.550 - 1.590). This indicates that a uniform and equal number (18,000 cells/well) of cells are being loaded into every well in the 96-well plate during the day-to-day operation. The cells reduced MTT to formazan giving maximum blue color uniformly and consistently.

YF-Control Compound-Virus Load Performance: A bar graph scatter plot of the mean virus load O.D. readings of the 42 control assays is presented in Figure 16-B. The results indicate that the average virus load O.D. reading is 0.240 (SD \pm 0.110) with a median of 0.240 (range of 0.030 - 0.480). This demonstrates that a good cell destruction is taking place and a uniform load of virus (32 TCID₅₀) is administered on the cell monolayer with very consistent viral CPE results.

YF-Control Compound-Assay Differential Performance: A bar graph scatter plot of the mean O.D. differential values of the 42 control assays is provided in Figure 17-B. The results indicate that the average differential O.D. reading is 0.870 (SD \pm 0.150) with a median of 0.872 (range 0.515 - 1.182). The single bell-shaped curve is reasonably sharp and uniform. This reflects that the assays are executed consistently and are repeatable during day-to-day operation with close to 87% measurement accuracy.





4.1.3.2 YF-Antiviral Activity Results:

Drugs with 95% Antiviral Reduction Levels: Out of the 6252 actual single drug tests, 62 new compounds demonstrated excellent antiviral activity, having antiviral reduction values of equal to or better than 95%. This represents around 1.0% of the test compounds being active at this excellent reduction level. These compounds are summarized in Table 7 according to the highest Total Antiviral Index (TAI). Compounds AVS-3102, 2631 and 5580 demonstrated the greatest in vitro promise, having TAI is of 99, 75 and 70% and Selectivity Indices (SI) of > 313, 98 and 76, respectively. The next nine compounds, demonstrated good antiviral activity with TAI's greater than 40% and SI values that ranged from 8 - 58. Twenty-nine other compounds demonstrated moderate antiviral activity, having TAI's from 25 -37% and SI's from 4.8 - 9.9. The rest (29 compounds) had only marginal antiviral activity with TAI's ranging from 5 - 24% and SI's of < 1 to 26.

It is worthwhile to note that compounds received in shipment number 62 were mostly colored (Table 7). Therefore those compounds appearing in the 95% active category from shipment number 62 should be interpreted with caution, since colored compounds create false positive readings with the MTT assay.

Table 7

AVS Compounds Active Against Yellow Fever Virus (YF) at AI₉₅ Level

Virus	AVS	Ship- ment#	Test	Diff- rntl.	IC 95		TC 95		NT 05	SI		ma T	
VILUS	NO.	Mench	Date	riici.	10 93		10 93		AI 95	31		TAI	
YF	3102	28	05/19/89		0.39	>	100.00	>	258.00 >	313.00	>	99.41	
YF	2631	65	07/06/90	1.083	9.30	>	1000.00	>	107.58	98.03			
YF	5580	54	05/03/89	1.295	6220.00	>	100000	>	16.10 >	75.60	>	70.14	
YF	3802	67	05/31/90		0.30		10.00	>	33.27 >	58.40	>	56.12	
	1019	28	03/06/90	1.032	6.22		100.00	>	16.08	30.74	>	55.70	
	4855	48	09/12/89		29.30		320.00		10.91 >	23.82			
	6225	62	01/16/90	0.996	30.10		320.00	>	10.62 >	18.33			
	2604	19	09/06/89		18.90		100.00		5.30	25.16			
	6218	62	01/16/90		44.30		320.00		7.22 >	17.12			
	0148	66	05/30/90		0.91		3.20		3.51 >	8.14			
	2591	19	09/06/89		30.20		320.00		10.60 >	17.89			
	2589	65	04/12/90		91.70		320.00		3.49 >	7.59			
	5121	56	11/01/90		97.90		320.00		3.27 >	8.30			
	6214	62	11/01/90				1000.00		10.59	8.96		35.36	
	8374	76	01/17/91				3200.00		3.44	8.12			
	2590	19	09/06/89		92.70	>	320.00	>	3.45 >	6.84			
	7067	72	10/03/90		28.30		906.00		31.97	9.86		33.97	
	0206	67	05/30/90	1.076			1000.00		10.77	9.72			
	6200	62	02/01/90		97.30		320.00		3.29 >	6.15			
	6201	62	02/01/90		93.80		320.00		3.41 >	6.07			
	2318	67	05/31/90		0.97		10.00		10.34	6.23		31.89	
	6219	62	11/01/90		94.20		320.00		3.40 >	5.83			
	6943	69	07/25/90		94.20		320.00		3.40 >	5.83			
	4113	39	09/11/89		0.57		1.00		1.77 >	5.96			
	2572 2980	65	04/12/90		93.30		320.00		3.43	5.80			
		61	12/06/89		0.90		3.20		3.56	6.03			
	6208	62	01/16/90		246.00		320.00		1.30 >	4.82			
	5138 6946	57 69	08/15/89		279.00		1000.00 966.00	-	3.53 3.47	5.90 5.73		27.36	
	2506	21	09/06/90 09/06/89		3.20		10.00	_	3.13 >	6.11			
	6220	62	01/16/90		93.70		320.00		3.42 >	6.16			
	6217	62	01/16/90		94.50		320.00		3.39 >	5.66			
	4281	61	12/06/89		2.98		32.00		10.75	5.45		25.99	
	8364	76	01/17/91	0.540	286.00		966.00		3.38	4.79		24.40	
	6195	62	02/01/90		267.00	>	320.00	>	1.20 >	4.33			
	6942	69	09/06/90				1000.00		1.40	3.12			
	6369	63	03/20/90		29.50	•	96.60		3.28	3.47			
	6460	63	03/22/90		93.20		874.00		9.38	3.54			
	7092	72	10/25/90			>	1000.00	>	1.02 >	2.11			
	6203	62	02/01/90		94.50		309.00		3.27	2.74		19.96	
YF	7068	72	10/03/90		30.80		293.00		9.51	3.83		19.91	
YF	6196	62	02/01/90		286.00	>	320.00	>	1.12 >	3.06	>	19.66	
	8563	76	01/03/91		93.80		320.00		3.41	3.94	>	19.43	
YF	2570	19	09/06/89	0.718	31.00		32.00		1.03 >	2.16			
	6197	62	02/01/90		296.00	>	320.00		1.08 >	2.19			
YF	2586	19	09/06/89	0.775	306.00		320.00		1.05 >	2.08	>	17.27	
YF	5405	66	07/06/90	1.155	0.31		8.19		26.12	3.67		17.25	
	6183	62	01/31/90		294.00	>	320.00	>	1.09 >	2.35			
	6185	62	01/31/90		94.10		315.00		3.35	2.89		16.47	
	4992	61	12/06/89		302.00		966.00		3.20	2.74		15.08	
	6207	62	10/31/90		302.00		966.00		3.20	2.74		15.08	
	5780	59	01/23/90		94.50		318.00		3.36	2.78		15.02	
	6186	62	02/01/90		96.60		313.00		3.24	2.73		14.62	
	2453	18	09/06/89		10.00		31.50		3.15	2.69		14.12	
	6205	62	02/01/90		299.00		320.00		1.07 >	2.00			
	2582	19	09/06/89		93.60		100.00		1.07 >	1.94			
	6227	62	01/16/90		302.00		320.00		1.06 >	1.79			
YF	2960	27	10/04/90	1.046	903.00	>	1000.00	>	1.11	0.00	>	9.92	

Table 7 (Cont'd)

Virus	AVS No.	Ship- ment#		Diff- rntl.	IC 95	TC 95	AI 95	sı	TAI
YF	7049	69	09/13/90	1.039	945.00 >	1000.00 >	1.06 >	1.77 >	9.27
YF	4074	48	12/07/89	1.211 <	1.00	26.80 >	26.82 >	1.60 >	6.33
YF	0361	2	CS/22/89	1.020 <	0.03	0.29 >	9.02 >	1.63 >	5.46
YF	6194	62	02/01/90	0.714	302.00	96.10	0.32	0.23 >	5.04

Drugs with 50% Antiviral Reduction Levels: Out of the 6252 actual single drug tests, 175 new compounds demonstrated good antiviral activity, having antiviral reduction values equal to or better than 50%. This represents around 2.8% of the test compounds being active at this good antiviral reduction levels. These compounds are summarized in Table 8 according to the highest Total Antiviral Index (TAI). AVS-4611 demonstrated the best TAI of 66% and SI of 109. Twenty-seven other compounds demonstrated moderate antiviral activity, having TAI's from 25 - 43 and SI's from 1 - 56. The rest (147 compounds) showed marginal antiviral activity with TAI's from <1 to 24% and SI's from <1 to 7.

It is worthwhile to note (Table 8) that compounds received in shipment number 62 were mostly colored. Therefore those compounds appearing in the 50% active category from shipment number 62 should be interpreted with caution, since colored compounds create false positive readings with the MTT assay.

Table 8 ${\tt AVS~Compounds~Active~Against~Yellow~Fever~Virus~(YF)~at~AI_{50}~Level}$

	AVS	Ship-	Test	Diff-										
Virus	No.	ment#	Date	rnt1.		IC 50		TC 50		AI 50		SI		TAI
YF	4611	65	07/06/90	0.874		6.54	>	1000.00	>	152.94		108.82	>	65.62
YF	2978	27	04/17/89			42.30	>	320.00	>	7.56		7.56	>	42.81
	0084	67	05/30/90			17.50		316.00		18.06		11.76		
YF	6618	64	04/19/90				>	1000.00	>	6.79	>	6.79		
YF	2503	67	07/12/90			0.58		10.00		17.13		17.13		
	6444	63	03/22/90			29.80		731.00		24.55		13.77		
YF	2573	65	04/12/90				>	1000.00	>	17.11		9.35		
	7438	73	09/20/90			1.94		206.00		106.01		55.49	> :	33.12
	4796	61	12/06/89			21.50	>	320.00	>	14.89		9.64		
YF	5998	61	02/01/90			2.01		32.00	>	15.92		8.34	> :	30.66
YF	8370	76	01/17/91	0.603		168.00		1920.00		11.46		7.64	> :	30.66
YF	0053	64	03/08/90			246.00	>	320.00	>	1.30		1.30		
YF	5450	65	04/26/90	0.708		2.68		65.30		24.42		11.96	> :	30.16
YF	4035	65	07/06/90	0.783		109.00	>	1000.00	>	9.17	>	9.17		
	4592	61	12/06/89	0.795		7.04		83.90		11.92		7.53		
	6209	62	01/16/90	0.912		55.00	>	320.00	>	5.82		5.82		
	8661	76	01/17/91			1.69		320.00		.189.07		13.00		
	2575	57	12/06/90			19.40		679.00		34.93		4.95		28.46
	6458	63	03/22/90					1000.00		4.27		4.27		
_	8511	76	12/20/90					3200.00		5.34		5.34		
	5484	66	05/08/90			76.70	>	320.00		4.17		4.17		
	8268	76	12/12/90		<	3.20		494.00		154.36	>	56.25		
	2600	65	04/12/90	0.843			>	1000.00	>	16.70		4.98		
	2563	21	09/06/89			0.17		2.20		13.17		6.00		
	4590	61	12/06/89			0.34		2.77		8.20		5.29		
	6223	62	01/16/90			27.70		216.00		7.78		5.69		25.20
	6707	67	07/12/90			12.10		95.00		7.82		5.23		
	4277	65	07/06/90			2.19		20.40		9.33		5.08		
	4609	63	02/13/90			0.03		0.26		8.31		4.23		24.39
	7087	72	08/31/90			3.88		31.50		8.10		4.65		24.20
	8261	76	12/12/90	0.826		170.00		2940.00		17.28		3.66		24.07
	8327	76	01/17/91				>	3200.00	>	1.35	>	1.35 5.52		
	2363 2309	15 53	08/29/89			8.18 34.70		64.50 297.00		7.88 8.56		5.26		23.69
	-		12/05/89					409.00				7.01		
	8377 3612	76 32	01/17/91 11/03/88			30.30		320.00		13.48		2.23		
	0360	2	08/22/89			0.05		0.29		6.23		4.19		22.90
	2581	65	04/12/90			52.40		320.00	-	6.10		4.99		
	2585	65	04/12/90			55.80		320.00		5.74		4.92		
	6202	62	10/31/90			154.00		660.00		4.28		3.17		22.15
	8378	76	01/17/91			148.00		868.00		5.86		4.01		
	5997	61	11/01/90			1.82		32.00		17.58		4.28		
	8332	76	01/17/91			177.00		1630.00		9.19		5.00		
	8262	76	12/12/90				>	1000.00	>	2.02	>	2.02		
	1381	45	02/13/90			13.60	-	87.70		6.45		4.40		20.78
	6234	67	11/01/90				>	3200.00	>	2.09	>	2.09		
	2812	61	12/06/89			0.01	•	0.04		8.66	Ť	4.75		
	4978	27	12/05/89			112.00		806.00		7.20		4.10		20.42
	4278	61	12/06/89			5.14		53.70		10.44		5.00		20.15
	6445	63	03/01/90			18.90		65.30		3.45		2.53		
	4527	63	02/13/90			3.66		26.90		7.36		5.05		19.83
	8225	75	12/06/90			32.00		202.00		6.30		0.86		
	2580	19	09/06/89			2.37	>	10.00	>	4.22	>	4.22		
	7085	72	08/31/90			6.31		56.30		8.92		3.49		19.39
YF	7383	70	09/27/90			158.00		680.00		4.29		3.03		
YF	6589	64	04/18/90				>	1000.00	>	5.01	>	5.01		
YF	4591	63	02/13/90		<	0.00		0.03	>	8.50	>	5.81	> :	19.16
YF	8371	76	01/17/91			431.00		2680.00		6.23		4.24	>	19.02
	7319	70	08/16/90			2.85		21.60		7.58		4.07		18.82
YF	7354	70	08/16/90	0.777		432.00	>	320.00	>	0.74	>	0.74	>	18.62

Table & (Cont'd)

	AVS	Ship-	Test	Diff-									
Virus	No.	ment#	Date	rnt1.	IC 50		TC 50		AI 50		SI	T	AI
	7904	74	10/24/90		6.34	>	320.00	>	50.48		1.27		.43
	2811	48	02/06/89		0.03		0.23		9.00		3.55		
	8353	75	11/30/90		58.60		320.00		5.46		2.78		
	8373	76	01/17/91			>	3200.00	>	3.71 >		3.71		
	6250	1P	11/14/89		88.50		286.00		3.24		2.18		
	2979	48	02/07/89		1.93		14.90		7.73		3.79		
	6979	68	06/21/90		88.50	>	320.00	>	3.62 >		3.62		.36
	6792	67	07/12/90		2.21		13.00		5.91		3.13		
	7083	72	08/31/90		21.90		147.00		6.69		2.37		
	0646	67	05/30/90		0.26	>	3.20	>	12.33		3.42		
	5483	66	07/18/90		2.88		0.24		5.34		3.68 3.96		.63
	4753 6256	44 1P	12/14/89 11/14/89		7.41		18.40 50.70		6.38 6.85		3.51		.55
	0094	1	11/01/90		239.00		742.00		3.10		1.68	_	
	3734	34	09/11/89		3.53		10.00		2.83).23		
	1644	64	11/01/90		17.90		76.10		4.25		2.49		
	1337	67	05/30/90	1.118	59.00		228.00		3.87		2.76		
	6456	64	04/18/90		61.00		198.00		3.25		2.16		
	7321	70	08/16/90		15.00		94.90		6.34		3.50		.50
	8372	76	01/17/91			>	3200.00	>	3.59 >		3.59		
	0083	64	03/08/90		1.28		9.16		7.14		1.82		
	6211	62	01/16/90		106.00	>	320.00	>	3.01 >		3.01		
	4223	63	02/13/90		0.22		0.90		4.08		2.77		. 58
YF	6477	66	07/11/90		18.00		76.20		4.22		3.00	14	. 54
YF	5916	60	10/23/89	0.874	8.85		46.90		5.30	- 2	2.73	> 14	.50
YF	5040	GABSN	02/08/90	0.848	268.00		2890.00		10.79		5.09	14	. 42
	5691	57	07/17/89		144.00	>		>	2.22 >		2.22		
	7424	70	09/27/90	0.932	320.00		595.00		1.86		1.23		.12
_	7065	72	10/25/90		18.30		81.80		4.46		2.20		
	8263	76	12/12/90		44.30		91.60		2.07		1.40		
7.1	7377	70	08/22/90			>	1000.00	>	1.05 >		1.05		
	7945	75	12/05/90		647.00		2110.00		3.27		2.41		
	5601		02/08/90		3720.00	>		>	0.86		0.73		
	7044	69	08/02/90		83.70		193.00		2.31		1.54		
	7084 6620	72	08/31/90		7.44 60.70		27.60 192.00		3.71 3.16		2.11		
	6986	64 68	04/19/90 06/21/90	0.363	63.60		255.00		4.02		2.74		.38
_	8269	76	12/12/90		100.00		940.00		9.40		1.40		
	4280	42	11/01/90		6.84		20.30		2.97		1.67		
	8271	76	12/12/90		189.00		830.00		4.40		2.52		
	6215	62	01/16/90		168.00	>	320.00	>	1.91 >		1.91		
	2579	19	09/06/89		214.00		320.00		1.50 >	_	1.50		
YF	7433	70	08/30/90		19.80		82.20		4.15		2.74		.62
	5539	56	08/07/89		2.27		6.08		2.68		1.31		
YF	7375	70	08/22/90		61.40		320.00		5.21		1.63		.51
YF	7332	70	08/16/90	0.902	100.00		207.00		2.07		1.28	11	.12
YF	2907	26	09/07/89	0.731	66.80		248.00		3.71		2.58	10	.97
	6226	62	01/16/90		184.00		320.00		1.74 >		1.74		
	6628	64	04/19/90		192.00	>	320.00	>	1.67 >		1.67		
	3935	65	07/06/90		84.30		210.00		2.49		1.84		.75
	5241	52	12/14/89		100.00		578.00		5.78		3.63		. 47
	4036	65	07/06/90		201.00		667.00		3.32		1.55		
	5186	58	10/03/89		74.90		341.00		4.56		2.63		.12
	4070	48	12/07/89		2.22		8.30		3.73		1.99		
	7071	72	10/25/90		226.00		495.00		2.19		1.01		
	8698	76	01/23/91		88.10		199.00		2.26		1.08		.87
	6417	66	07/06/90		86.40		396.00		4.58		1.84		.67
	6147	62	01/04/90		2.66		6.60		2.48		1.84		.63
	6626 9123	64 77	04/19/90 01/31/91		243.00 : 296.00		320.00		1.32 > 6.16		1.32		.52
	7910	75	11/08/90		1870.00	>		>	1.71 >		1.71		.16
	6228	62	01/16/90		222.00		320.00		1.44 >		1.44		. 15
1.5	3220	72	01/10/30	J. 302	222.00		320.00		1.44 /		44	- 7	. 13

Table 8 (Cont'd)

	AVS	Ship-	Test	Diff-										
Virus	_	ment#		rntl.		IC 50		TC 50		AI 50		SI		TAI
VILUE	110.	merica	Date	Incl.		10 30		10 30		WT 20		31		1112
VE	2274	12	08/29/89	0.733		65.20	>	100.00	>	1.53	>	1.53	>	8.90
	6212	62	01/16/90			206.00				1.56		1.56		8.66
	6044	61	02/08/90			0.10		0.66		6.60		4.90		8.63
	6315	63	03/20/90		Ť	7.62		20.50		2.69		1.94		8.59
	7378	70	08/22/90			257.00		614.00		2.38		1.63		8.45
	6617	64	04/19/90					1000.00	>		>	1.60	>	8.43
	2992	27	09/07/89			81.60						1.23		8.39
	6137	62	12/28/89			9.31		21.00		2.26		1.66		8.29
YF	7086	72	08/31/90			6.40		10.00		1.56		1.03	>	8.07
YF	4739	44	07/06/89			29.00		67.60		2.33		1.65		8.03
	4452	44	07/06/89			78.50		301.00		3.84		1.84		7.92
	5781	59	01/30/90			195.00		601.00		3.08		0.44		7.87
	5543	56	10/04/90			28.10		77.90		2.77		1.36		7.77
	5995	61	11/07/89			8.19		22.20		2.71		1.08		7.66
	5535	56	06/13/89			185.00	>		>	1.73	>	1.73	>	7.36
	7472	73	10/31/90			100.00		213.00		2.13		0.85		7.34
	3980	36	09/11/89			65.70		100.00	>	1.52	>	1.52		7.22
	6334	63	03/20/90			3.09		11.60		3.77		1.75		7.11
	7032	69	08/01/90			230.00						1.39		6.98
	5538	56	06/13/89			215.00			>	1.49	>	1.49	>	6.96
	1736	45	12/07/89			134.00		547.00		4.09		0.83		6.78
	5197	58	10/04/90			86.70		159.00		1.83 2.28		0.98		6.76
	6326 6422	63 66	03/20/90 07/06/90			8.64 313.00		19.70 657.00		2.10		1.20		6.71
	8240	75	12/06/90			714.00		2230.00		3.13		1.59		6.17
	3592	61	12/06/89	0.813				1000.00		1.24	,	1.24		6.06
	8326	76	01/17/91			259.00		568.00		2.19		1.30		5.80
	7390	70	08/22/90			245.00		558.00		2.28		1.38		5.78
	2902	26	09/07/89			65.20			>		>	1.53		5.72
	9128	77	01/31/91					3200.00		1.07		0.17		5.41
	4768	44	11/08/88			74.90		173.00		2.31		1.34		5.40
	7445	73	10/25/90				>	3200.00	>	1.25	>	1.25		4.64
	1976	1	08/29/89			80.40				1.24	>	1.24		4.32
YF	6683	64	04/05/90			204.00	>	320.00	>	1.57		1.35		4.25
YF	5072	48	02/27/89			230.00	>	320.00	>	1.39		0.38	>	4.01
YF	8395	76	12/19/90			752.00		1450.00		1.93		0.79		3.92
YF	1838	64	11/01/90	0.951		252.00		558.00		2.22		1.34		3.75
YF	7003	69	09/06/90	1.009		252.00		556.00		2.21		1.31		3.75
	7434	70	10/03/90			264.00		570.00		2.16		1.34		3.65
	6199	62	02/01/90	0.712		245.00			>	1.31		0.90		3.49
	7373	70	08/22/90			252.00	>		>	1.27		1.20		3.35
	4785	46	11/01/90			3.20		8.09		2.53		0.89		3.20
	5905	61	10/30/89	0.706		8.85		16.10		1.81		0.94	>	3.03
	4754	44	12/20/89			7.99		11.20		1.41		0.85		2.08
	7045	69	08/02/90			320.00				1.00		0.37		1.67
	7051	72	10/03/90				>	1000.00	>	1.32		0.54		1.02
	8311 6309	76 63	12/12/90 03/20/90			864.00 95.10		1240.00 250.00		1.43		0.33	,	0.71
	7461	73	10/31/90			84.10		29.50		0.35		0.38		0.59
	8404	75 76	10/31/90			896.00		959.00		1.07		0.50		0.47
	8693	76	01/23/91			320.00				1.00		0.07		0.47
	6204	62	02/01/90			30.00	-	22.80		0.76		0.55		0.00
	8251	76	12/12/90			320.00		217.00		0.68		0.50		0.00
	2231	, 0	12/12/30	3.307		223.00		217.00		0.00		5.50		3.00

<u>Drugs with 25% Antiviral Reduction Levels:</u> Of the 6252 actual single drug tests, 394 new compounds demonstrated moderate antiviral activity, having antiviral reduction values equal to or better than 25%. This represents around 6.3% of the test compounds being active at this marginal antiviral reduction level.

In general, when compared to the 95% and 50% antiviral activity categories, the compounds in this (25%) category do not appear to have any significant antiviral promise and probably do not need to be presently confirmed any further.

4.1.3.3 Confirmatory Assays:

Some of the compounds were sent to us in more than one separate drug shipment. These compounds were tested more than once. Data from the confirmatory assays are summarized in Table 9. If a compound showed $\geq 50\%$ reduction in CPE during this contract period, then it was considered a candidate for confirmatory testing. The confirmatory tests are from active compounds picked by both the VR and MTT assays. Out of 178 confirmatory tests, 137 compounds were confirmed active during this reporting period and the remaining 41 compounds gave conflicting results. The criteria for activity is that the confirmatory test has to show $\geq 25\%$ reduction in CPE. Failure to confirm the activity in these compounds was probably due to differences during the assay conditions:

- 1) In confirmatory assays the concentration range is adjusted to a more accurate semilog scale to maximize the TAI window.
- 2) Differences in the "differential" of the two runs can cause the compound to read positive or negative, falsely. The variability in the differential can cause false positive or negative bias to be introduced into the calculations, thus reflecting the variability in the maximum activity of the compound.
- 3) The metabolic rate, cell and virus load/well, age, and passage number of the cells may cause the above observed variability in the confirmatory results.
- 4) Problems associated with stability and storage of the compound (i.e., different lot numbers, solubility, light sensitivity, hygroscopic, etc.).
- 5) Problems associated with technical execution of large numbers of plates by different technicians.

During this reporting period the overall confirmatory rate against YF was 77%. The conflicting results should be retested at a later date based on the availability of the compound.

4.1.3.4 Recommendations of YF-Actives Based Upon the In Vitro Results with MTT Assay (Vero Cells).

Based upon the in vitro results with the MTT assay (Vero cells) we recommend the following:

- 1) Compounds with the highest TAI in the 95% activity category that have confirmed results with the exception of "colored" compounds should receive the highest priority for further profile studies and in vivo animal testing.
- 2) Compounds with the highest TAI in the 50% activity category that have confirmed results with the exception of "colored" compounds should receive the next highest priority for further profile studies and in vivo animal testing.

aple 9

Confirmatory Assays for Compounds Active Against Yellow Fever Virus

< U +		* * *	• • •	• • •	+ •	+ + +	• • •	* * • * * * *	+ + 1	+ + + + + +
ASSAY			### ####		CPE	9 9 9 9 9 9		222FFFF	CPE CPE	CPE CPE MIT MIT
TAI	30.54 1.23 0.01	15.11 9.67 9.14	0.32 10.57 37.83	9.21 1.30 16.29	0.00	1.40	25.62 42.52 3.05	1.40° 0.60° 0.50° 2.22 4.73 13.21	2.60 1.10 0.73	1.60° 0.00° 11.70 0.41 22.90 10.18
15	1.30	4.82 > 1.39 > 0.00 >	1.40 >	0.00 0.	2.10	0.20	4.38 × 8.14 × 0.70	0.50 0.00 0.00 0.00 0.00 0.00 0.00 0.00	10.00 0.003 0.00	6.60 0.00 0.00 4.19 2.23
A1 95	888	988	9.00	0.00	0.00	00.00	3.51 >	0.01 0.00 0.00 0.00 0.00 77.01	0.00	3.20
70 95	320.00 279.00 320.00	30.40 9.92 28.30	320.00 100.00 970.00	100.00 1000.00 2850.00	320.00	0.32 10.00 10.00	3.20	320.00 - 320.00 - 320.00 - 1000.00 -	320.00 320.00 320.00	0.32 v 10.00 2.93 1.00 1.00
10 %	98.9	0.00	0.00	0.00	320.00 > 0.00 >	0.00	0.00	320.00 × 0.00 ×	0.00	0.00000
AI 50	0.00	7.14 2.00 0.00	1.40	0.00 3.10	0.00	0.20	55.79 8.14 2.93	2.80 ~ 1.10 ~ 1.50 0.00 0.00 0.00 16.57	32.00 3.20 > 0.00	6.60 - 0.00 6.47 6.23 3.13
10 50	320.00 > 9.86 87.60	9.16 6.40 7.73	73.30 100.00 > 316.00	100.00 694.00 742.00	320.00 > 235.00	1.00 0.32 0.10	3.20 > 2.52	320.00 > 320.00 > 320.00 > 320.00 > 320.00 > 320.00 320.00 320.00 742.00 791	320.00 320.00 > 320.00	0.32 v 10.00 0.51 0.00 1.00 v
10 50	246.00 × 0.00 0.00	1.28 3.20 0.00	0.00 71.50 > 17.50	0.00 > 0.00 239.00	48.00 >	0.48 1.00 0.00	0.06 0.39 v	115.00 > 297.00 > 217.00 > 0.00 > 0.00	9.90 100.00 v	0.07 0.00 0.00 0.00 0.03 0.03
AI 25	320.00 0.00 0.00	6.18 3.67 3.47	3.29	9.83	3.20	1.00	10.03 18.28 1.48	1.00 0.00 1.68 5.11 7.44	1.00	10.00 0.00 0.00 7.39 5.25
70 25	320.00 > 6.46 22.90	6.18 > 4.45 5.47	42.90 100.00 > 206.00	100.00 279.00 402.00	100.00 -	0.07	3.20 \$	320.00 - 100.00 - 100.00 - 459.00 + 464.00	100.00 ~ 0.32 88.00	0.32 ~ 10.00 0.21 0.20 0.71
10 25		1.21	30.40 >	0.00 0.00 40.80	32.00	5.20 3.20	0.03	- 320.00 - 320.00 0.00 191.00 > 89.70 53.90	0.32	0.032 0.00 \$ 0.05 0.03 0.14
Diff.	0.985	0.973 < 1.030 0.984	1.252 0.897 1.062	0.793	1.088	\$\$\$	1.076	NA NA NA NA NA NA NA NA 0.705 0.739 1.076	NA NA 0.986	NA 1.193 1.193 0.842 0.915
=	UF2 VSS 280	UF4 VSU ZXA	457 40A	\$25 767 2X8	: g	:::	28 P P P P P P P P P P P P P P P P P P P	1 : : 8 8 8 9 1 : : 8 8 8 8 9 9 9 9 9 9 9 9 9 9 9 9 9 9	:::	SXB
Test	03/08/90 05/01/90 10/04/90	03/08/90 05/01/90 11/01/90	05/22/89 08/22/89 05/30/90	10/10/89 04/10/90 11/01/90	07/06/88 05/08/89	02/04/88 05/04/88 05/18/88	08/22/89 05/30/90 10/04/90	01/30/87 02/02/87 05/24/88 05/08/89 12/05/89 12/05/89 05/30/90	01/15/87 08/19/87 01/23/89	03/06/87 05/18/88 02/06/89 02/06/89 08/22/89
Ship-	2222 7	333	1-19	1 65	7 7		2,67	2,2,2,2,2,0	3	774477
AVS.	** VIRUS 0053 64 0053 64 0053 64	0063 0083	7800	7600	1110	0100	0148 0148 0148	9050 9050 9050 9050 9050 9050 9050 9050	0303 0303 0303	0360 0360 0360 0360

Table 9 (Cont'd)

Confirmatory Assays for Compounds Active Against Yellow Fever Virus

∢ ∪⊢	+ + + + +	+ + + + + +	++ ++++		* * * * * * * *
ASSAY TYPE	SEEFE	9 9 9 E E	S S S E E E		FFF FFF 99
TA1	1.10 14.54 5.46 22.59 15.48	3.10 0.60 1.70 3.20 9.18	3.00° 1.30° 3.50° 1.10° 10.00 5.36 55.70	0.00 11.66 13.23 4.68 0.00 15.56	23.99 7.87 15.73 2.27 0.08 3.75 2.30 1.60
ıs	2.7 2.72 1.63 × 5.29 2.76 ×	13.00 0.10 1.20 0.70 2.10 3.42 >	13.00 0.60 1.00 0.00 ~	0.00 2.38 1.38 > 0.00 2.76 > 4.40 >	0.00 \ 2.49 \ \ 0.92 \ \ 0.00 \ 1.34 \ 1.60 \ 1.60
AI 95	3.20	00.00 1.00 200.00 0.00 0.00	1000.00 1.00 1.00 1.00 16.08	, 8888888 888888 888888 88888 88888 88888	0000 0000 000
26 21	0.32 ~ 0.28 ~ 0.29 ~ 0.32	320.00 >1000.00 1.00 > 1.00 320.00 >1000.00 320.00 >3200.00 3.20 0.00	320.00 320.00 1.00 > 10.00 1.00 > 10.00	32.00 320.00 320.00 1000.00 1200.00 806.00 309.00	320.00 1000.00 869.00 320.00 943.00 956.00 320.00
10 98	0.00	0.32 × 0.10 × 0.10 × 0.10 × 0.10 × 0.00 × 0.00 ×	0.00 0.32 1.00 v 0.00 v 6.22 v	A A A A A	0.00 0.00 0.00 0.00 0.00 320.00
AI 50	8.00 4.41 2.35 <- 7.74 3.99	13.00 0.10 1.20 0.70 7.60 12.33		0.00 3.57 1.38 0.00 3.87 2.41	0.00 0.00 1.23 0.00 2.22 13.20
1C 50	0.32 - 0.07 > 0.08 0.08	3.20 0.032 0.10 0.032 2.17 3.20 >	32.00 13.00 10.00 0.60 320.00 -1000.00 > 1.00 > 3.80 ~ 2.35 0.00 1.00 > 108.48	32.00 228.00 320.00 > 286.00 320.00 228.00 87.70	320.00 716.00 76.10 320.00 × 433.00 558.00 320.00
10 50	0.06 0.03 0.01 0.02	0.20 0.50 0.08 0.04 0.29	2.50 17.40 0.32 ~ 0.00 0.00 ~	232.00 v 0.00 v 0.00 v 59.00 v	0.00 > 0.00 > 0.00 > 0.00 > 0.00 > 0.00 > 0.00 > 0.00 > 0.00
AI 25	3.20 4.76 1.63 < 12.43 4.83	10.00 32.00 1.00 0.66 3.68 5.50	32.00 1.00 1.00 < 1.00 0.71 2.18 60.56		26.06 0.00 8.10 1.63 2.13 67.00
55 21	0.10 0.05 0.06 0.06 0.05	2.10 0.021 0.021 0.60 0.89	21.00 6.60 0.32 ~ 1 0.10 ~ 1.00 >	9.50 320.00 × 191.00 × 320.00 × 163.00	240.00 × 44.60 × 44.60 × 240.00 × 126.00 × 210.00 × 210.00 × 21.00
10 25	0.032 0.01 0.01	0.21 0.66 0.03 0.032 0.16	6.60 6.60 0.32 0.10 0.45 0.47	A A	12.30 > 0.00
Diff.	1.182 1.020 < 0.944 0.865	NA NA NA 1.020 1.079	NA NA NA * 1.130 0.968 1.032	1.064 0.897 0.847 0.945 0.996 1.118 0.888	0.976 1.138 0.865 0.976 0.997 0.951
==	2XC 2XC	::::\$: : : : : : : : : : : : : : : : : : :	RB1 U19 VVZ VVZ WDE U11	2XC 2XC 2XC 3XC 3XC 3XC 3XC 3XC 3XC 3XC 3XC 3XC 3
Test Date	03/01/88 02/06/89 08/22/89 02/13/90 11/01/90	03/06/87 08/19/87 12/14/87 05/17/88 08/22/89 05/30/90	02/02/87 07/29/87 11/30/87 02/02/88 05/08/89 001 08/22/89 RAS 03/06/90 UD0	08/28/89 01/18/90 03/13/90 04/12/90 05/03/90 05/30/90 05/30/90	03/13/90 05/03/90 11/01/90 03/13/90 05/03/90 11/01/90 01/16/87
Ship-	~ 8 ~ 8 ~	200000	25 25 25 25 25 25 25 25 25 25 25 25 25 2	E24244 22	333 333
AVS.	0361 0361 0361 0361	97799	0703 0703 1019 1019 1019		1644 1644 1644 1638 1838 1838 1975
	00000	000000	00	eeeeee ee	++

Table 9 (Cont'd)

Confirmatory Assays for Compounds Active Against Yellow Fever Virus

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ASSAY	9 8		CPE .	TH	H	. GE	GE	CPE	CPE	20	S	HTT	MT	CPE		S	CPE	CP	8	CPE	TH		T T	SP	TH			H	ĔĒ	
Ţ	2.90		0.80	0.0	4.32	1.90	0.00	3.20	0.20	3.80	2.70	8.90	0.25	3.33	3.20	0.50	0.50	2.50	0.20	1.00	1.20	20.80	12.98	3.20	0.40	14.62	28.34	23.84	6.50	
IS	17.70	000	1.80	0,00	1.24 >	10.20	0.00	45.90	0.00	26.30	5.60	1.53 >	0.0	24.30	24.40	0.10	0.13	36.70	0.0	3.90	0.60	3.42 >	0.00	2.00	0.00	0.50	7.02 >	5.52 >	0.0 0.0	
N %	3.20	00.0	0.00	0.00	0.00	1.00	1.00	1.00	0.00	10.00	32.00	0.00	0.00	10.00	3.20	9.0	0.00	0.00	0.00	0.00	0.00	0.0	88	100.00	0.00	0.0	 	0.00	0.0	
10 95	320.00 >	100.00	320.00	320.00	100.00	320.00	320.00	320.00 >	320.00	320.00	320.00 >	100.00	320.00	320.00 >	320.00 >	320.00	320.00	320.00	320.00	320.00	320.00	320.00	986.00	320.00 >	320.00	10.00	0.00 0.00 0.00	99.20	282.00 287.00	
10 95	100.00	0.00	0.0	0.00	0.00	320.00	320.00	320.00 >	0.00	32.00	10.00 >	0.00	0.00	32.00 >	100.001	0.00	0.00	0.00	0.00	0.00	0.00	0.00	88	3.20 >	0.00	• 0.0 • 0.0	0.00	0.00	 0.0	
AI 50	07.71	800	9.	0.00	1.24	10.20	0.00	45.90	0.00	26.30	5.60	1.53	0.00	24.00	24.40	0.10	4.00	36.38	0.00	3.90	9.00	3.42	0.00	200.00	0.00	19.12	25.07	7.88	9.0	
10 50	320.00	100.00	100.00	183.00	100.00	320.00	3.20	320.00	32.00	32.00	10.00	100.001	203.00	320.00 >	320.00	10.00	320.00 -	100.00	32.00	320.00	320.00	320.00 >	297.00 671.00	320.00 >	2.51	10.00	10.00	64.50	20.70	
10 50	18.00	800	57.00	0.00	80.40 >	31.50	0.00	7.40	0.00	1.20	3.90	65.20 >	0.00	13.10 >	13.10	09.62	79.50 ~	2.70	0.00	\$1.90 >	53.20 >	93.70 >	34.79 0.00	1.60 >	0.00	0.52 >	0.40	8.18	0.0	
AI 25	32.00	1000.001	3.10	0.00	2.28	32.00	0.01	3.20	0.00	21.00	2.10	2.07	0.00	32.00	21.00	0.21	0.30	9.0	0.00	3.90	1.00	7.71	12.38 6.58	3.20	1.00	0.97	15.00	12.76	6.15	
52 21	210.00	100.00	90.99	93.50	100.00	2.00	2.10	210.00	21.00	21.00	9.60	100.001	135.00	320.00 ~	210.00	9.60	10.00	99.00	21.00	320.00	32.00	320.00 >	183.00 495.00	3.20 ~	1.00	0.56	2.83	45.10	55.90 13.30	
52 21	9.60	0.10	21.00	0.00	43.90 >	9.60	210.00	99.99	0.00	1.00	3.20	19.70 >	0.00	10.00	10.00	32.00	32.00	100.00	0.00	81.90	32.00	41.50 >	75.20	1.00		0.27	0.75	3.54	0.00	
Diff.	\$:	× * *	ž	0.680	0.851	¥	¥	×	¥	××	××	0.733	1.069	NA -	¥	¥	NA .	*	X	1 ¥	¥	0.955	0.765	¥	0.955 <	0.872	0.666	999.0	1.046	
ž*	; ;	:	:	916	*OX	:	;	:	:	:	:	ROP	169	;	:	:	:	;	:	:	:	PER	SVH				X FS	RDR	33	
Test	01/16/87	12/14/87	05/18/88	05/15/89	08/53/89	01/20/87	08/19/87	01/23/87	08/19/87	01/23/87	07/29/87	08/29/89 RDP	05/31/90 WG9	01/19/87	02/19/87	08/19/87	08/19/87	02/13/87	08/19/87	02/13/87	05/25/88	04/10/89 PER	12/05/89	03/04/88	04/10/89	08/55/80	05/51/90	08/55/89	03/13/90	
Ship- ment			_	22	_	_	_	2	2	2	~	12	29	12	12	12	12	=	=	=	=	23	22	13	53	٦	29	15	33	
No.	1976	1976	1976		1976	1978	1978	1983		1985	1985	2274		2275	2275	2275	2275	2305		2309			2303	2318	2318		2318 2318		2363	

Table 9 (Cont'd)

Confirmatory Assays for Compounds Active Against Yellow Fever Virus

∢ ∪⊢	• • •		• • •	+++++++	• • • •	+ + + +	* * * *	* *
ASSAY	EEE	CPE		888 ======			EE EE	FF
TĀĪ	14.12 2.85 0.02	3.60° 0.50 32.81 35.26 5.89	26.99 31.15 4.01	2.30 1.60 1.60 1.60 25.60 25.60 14.35	19.20 13.53 21.02 33.37	28.45 29.53 10.25	11.89 23.48 17.27 14.96	35.03
15	2.69	1.80 0.00 v 12.20 8.25 17.13 v	6.11 > 5.65 > 0.00 >	5.10 2.40 3.47 0.00 0.00 2.81 4	2.49	4.35 0.00 0.00 0.00	1.94 × 4.48 2.08 × 1.87 ×	6.84 v 9.96
¥ 8	3.15	320.00 0.00 0.00 0.00 0.00	3.13 > 0.00	6.0.0.0.0.0.0 0.0.0.0.0.0.0 0.0.0.0.0.0.	0.00 0.00 0.00 0.00 0.00	, , , , , , , , ,	1.07 \ 0.00 \ 0.0	3.45 > 10.35
10 %	31.50 31.80 29.90	320.00 × 320.00 10.00 10.00 10.00 32.00	10.00 × 320.00 320.00	10.00 3.20 3.20 3.20 0.00 10.00 10.00 10.00	32.00 × 311.00 320.00 1000.00	10.00 10.00 97.00 94.00	100.00 × 1000.00 × 320.00 × 320.00	320.00 >
S 21	0.00	- 00000 88888 88888	3.20 > 0.00 > 0.00 >	0.0000000000000000000000000000000000000	31.00	0000	93.60 v 0.00 v 306.00 v v 0.00 v	92.70 > 96.60 >
AI 50	3.66	571.00 ~ 0.00 45.48 19.44 17.13 4.22	6.11 18.35 0.00	31.00 - 5.10 - 25.00 - 6.07 0.00 4.90 13.17 0.00 5.06 5.06		34.93	1.94 14.01 2.08 1.87	6.84
10 50	21.30 21.50 10.80	320.00 × 2.52 10.00 × 10.00 × 10.00 × 3.02	10.00 > 94.10 31.40	10.00 3.20 2.20 10.90 1.00 1.00	32.00 × 320.00 × 1000.00 ×	679.00 10.00 × 63.40 40.40	100.00 × 1000.00 × 320.00 × 320.00 ×	320.00 > 917.00
10 50	5.83 0.30 0.00	0.56 v 0.00 0.22 v 0.51 v 0.58 v	1.64 v 5.13 0.00	0.32 0.63 0.13 0.00 0.00 0.017 0.00 0.00	14.80 × 21.40 × 68.80 × 58.50 ×	2.37 2 0.00	51.60 × 71.40 × 153.00 × 172.00 ×	46.80 > 57.20
A1 25	3.63	1.00 1.00 19.50 13.27 25.44 2.09	8.88 28.97 0.63	10.00 3.20 6.62 0.00 9.32 4.86	4.25 4.52 6.88 12.98	7.85 11.35 0.00 8.32	2.70 6.70 3.06 2.84	13.31
10 25	15.70 15.70 6.19	1.00 2.68 4.25 10.00 0.83	10.00 × 29.00 × 16.50	3.20 3.20 3.20 3.20 3.20 5.20 5.20		96.20 10.00 v 44.70 22.80	100.00 × 320.00 × 320.00 ×	320.00 > 570.00
10 25	4.32 9.19 0.00	1.00 < 1.00 < 0.32 0.32 0.40	1.13 > 26.40	0.32 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.10 < 0.	7.52 × 11.80	12.30 0.88 > 0.00 2.74	37.00 × 47.80 105.00 × 113.00 ×	32.00 >
Diff.	0.736	NA - 0.986 > 0.724 1.121 0.526 1.074	0.715 1.068 < 1.011	NA N	1.101	0.980 0.820 0.752 0.980	0.73 0.952 0.73 0.958	0.619
# P.	/89 RFF /90 UID /90 WZ	/87 /89 PES /89 RFG /90 WGB /90 XFT	/89 RFH /90 UIE /90 W2	/87 /88 /89 0JW /89 0JW /89 0SB /89 RFI /89 SRG		/90 1S7 /90 VBL /90 1S7	/89 RFL /90 VBH /89 RFL /90 1SB	09/06/89 RFM 04/12/90 VBN
p- Test t Date	09/06/89 03/13/90 05/03/90	07/16/87 04/10/89 09/06/89 05/31/90 10/04/90	09/06/89 03/13/90 05/03/90	06/25/87 03/01/88 05/25/88 02/06/89 05/22/89 09/06/89 12/05/89	09/06/89 12/06/90 06/26/89 04/12/90	12/06/90 09/06/89 04/12/90 12/06/90	09/06/89 04/12/90 09/06/89 12/06/90	09/06
Ship-	223	21 23 67 67	222	21 32 48 48 48 15 21 15/21	57 59 57	52 65 65	5 2 5 5	65
AVS No.	2453 2453 2453	2503 2503 2503 2503 2503 2503 2503 2503	2506 2506 2506	2563 2563 2563 2563 2563 2563 2563 2563	552 553 553 553 553 553 553 553 553 553	25.75 25.86 25.80 25.80 25.80	2582 2582 2586 2586	2590

Table 9 (Cont'd)

Confirmatory Assays for Compounds Active Against Yellow Fever Virus

∢ ∪⊢	+ +	+ +	+ +		+ •	+ 1	+ + +	+ + +	** **		+ + + + +
ASSAY	FI	FF	CPE	FFF	CPE	H		CPE CPE			SELEE
IAI	39.96	46.09	3.80	63.24 49.47 75.11	1.20	18.20	12.11 20.52 0.00	2.30 2.10 4.85	10.97	0.48 42.81 0.00 16.57	1.00 17.57 8.93 11.98 0.00
S	17.89 > 4.17 >	25.16 >	49.80	58.39 > 19.50 > 98.03 >	1.30	3.55 > 0.00 >	4.73 \$	12.90	0.00	0.00 7.56 v 0.00 2.57 v	2.10 2.02 0.00 0.00
A1 95	10.60 >	5.30	3.20	12.06 > 4.32 > 107.58	0.00	0.0	0000	1.00 3.20 0.00	0.00	00000	3.20 0.00 0.00 0.00
26 21	320.00 >	100.00	320.00 >	320.00 > 320.00 > 1000.00 > 1	100.00 ~ 93.70	0.00	0.01	320.00 320.00 >	320.00 320.00 320.00	320.00 320.00 100.00	32.00 28.60 95.60 89.90 9.09
S 31	30.20 >	18.90 \$	100.00 > 32.00 >	26.50 > 74.10 > 9.30 >	0.00	0.00	0.00	~ 320.00 > ~ 100.00 > 0.00 >	0.00		0.0000
AI 50	17.39	29.56	49.80	58.39 19.50 150.71	0.00	0.00	3.28	13.00	0.00		6.80 3.73 0.00 0.00
TC 50	320.00 >	100.00	320.00 100.00	320.00 > 320.00 > 726.00	100.00 37.00	0.23	0.01	320.00 320.00 97.20	248.00 157.00 41.70	320.00 320.00 > 100.00	32.00 14.90 24.70 23.00 2.53
10 50	17.90 > 57.60 >	3.38 > 7.30 >	6.40	5.48 > 16.40 > 4.82	7.60 - 0.00	0.03	0.00	24.60 > 23.10 > 0.00	66.80 0.00 213.00	0.00 × 42.30 × 0.00 × 262.00 ×	4.70 1.93 6.55 0.00
AI 25	53.93	55.93	× 100.00 6.60	185.96 40.19 141.25	3.10	6.08	4.57 7.12 0.00	13.00 3.20 1.30	1.04	0.00 320.00 0.00 5.91	3.10 10.80 3.03 7.27 0.00
10 25	320.00	85.10	320.00	320.00 > 320.00 > 472.00	10.00	0.00	0.01	320.00 32.00 62.00	172.00 69.80 13.10	M W - 40	10.00 7.31 13.20 12.90 1.64
10 25	13.40 > 37.40	1.52	3.20 <	1.72 > 7.96 > 3.34	3.20	0.00	0.00	- 24.60 10.00 47.00	36.90 67.40 146.00	0.00 \$	3.20 0.68 4.36 1.77 0.00
Diff.	0.619	0.650	\$ \$	0.650 1.066 1.083	NA 0.962	1.18	1.135 0.813 1.033	NA 1.020	1.068		1.083 1.067 1.045 0.754
ž*	V80	D RFN	::	N XON	2	200 6	0 0K0	3 ::	RHC UIE		P023
Test	09/06/89	09/06/89	04/17/87	09/06/89 04/17/90 07/06/90	05/27/87	02/06/89	02/06/89 12/06/89 12/06/90	09/14/87 05/25/88 04/10/89	09/07/89 03/13/90 09/07/89	02/28/89 04/17/89 05/15/89 02/13/90	09/01/87 02/07/89 03/10/89 04/17/89
Ship- ment	\$ 5	5 5	5 5	6 5 5 5	75	84	825	22 52	22 22		22 48 52 24 25
AVS No.	2591 2591	2604	2630	2631 2631 2631	2716 2716	2811	2812 2812 2812	2906 2906 2906 2906	2907	2978 2978 2978 2978	2979 2797 2797 2797 2797

Table 9 (Cont'd)

Confirmatory Assays for Compounds Active Against Yellow Fever Virus

∢ ∪⊢	+ + + + + +	+ + + + +	• • • • •	+ + +	+ + + + +	** ***	
ASSAY	CPE CPE MITT MITT				MATT THE	CPE ATTE	
ŢĀ	2.30° 2.40° 32.11 10.03 13.49 28.76	3.53 0.00 1.48 6.06 0.00	55.42 55.42 56.12	0.00 0.50 10.75	1.00° 11.68 11.51 3.43 29.41	7.43 10.30 1.40 6.10 5.01	0.00 2.81 6.33 31.63
18	5.10 2.30 14.78 v 1.80 4.78 v 6.03 v		29.30 × 58.40 × 58.40 ×	0.00 0.00 1.84	1.70 0.00 2.00 9.17 ×	1.25 v 1.55 v 1.57 1.42 v 1.99 v	0.00
A1 95	10.00 1.00 0.00 0.00 3.56	000000	0.00 0.00 0.00 11.13 >	0.00	0.00	00 0000	0.00 0.00 26.82 v
5 8	32.00 10.00 3.20 3.20 10.00 3.20 v	320.00 320.00 1000.00 320.00	1000.00 1000.00 10.00.00 10.00 v	10.00 > 308.00 272.00 309.00	320.00 160.00 32.00 320.00	320.00 1000.00 10.00 3.20 3.20 66.00	0.03 0.03 26.80 v
S 21	3.20 0.00 0.00 0.00 0.90 0.90			^	0.00 0.00 0.00 0.00 0.00	00.000000000000000000000000000000000000	0.0000
AI 50	51.00 ~ 2.30 35.96 4.08 9.92	0.00	0.00 0.00 0.00 29.30 58.40	0.00 0.00 2.49	1.70 0.00 3.87 0.00 9.17	1.30 3.32 1.80 ~ 1.57 1.42 3.73	0.00 0.00 2.68 <
TC 50	32.00 3.20 2.35 2.53 2.17	320.00 320.00 320.00 1000.00 >	1000.00 1000.00 10.00 10.00 10.00	10.00 × 204.00 74.10 210.00	32.00 97.80 32.00 >	320.00 > 667.00 3.20 3.20 > 3.20 > 3.20 > 3.20 > 8.30 3.20 > 3.20	0.03 0.03 2.68 v
10 50	0.63 1.40 0.07 0.62 0.32		0.00 1 0.0	A	19.20 0.00 8.30 > 0.00 > 1	246.00 × 201.00 1.80 2.04 × 2.22 2.22	0.00 \$
AI 25	10.00 10.00 21.30 2.58 7.70 12.09			71.73 0.00 3.75	3.20 5.80 4.00 11.07	2.41 2.77 3.20 2.35 3.65	0.00 1.72 1.60 <
TC 25	3.20 3.20 0.97 1.12 2.10	320.00 320.00 320.00 7 1000.0 7 276.00	973.00 v 10.00 v 10.00 v	10.00 × 146.00 50.20 155.00	32.00 49.00 17.00 245.00	306.00 312.00 3.20 3.20 >	0.03 0.03 > 1.60 >
10 25	0.32 0.32 0.05 0.43 0.14	A A A		A	10.00 8.40 4.20 230.00 54.40 >	127.00 112.00 1.36 × 1.49 × 1.21	0.00 0.00 0.00 0.00 0.00
oiff.	NA NA 1.154 0.971 0.676		1.055 0.760 1.017 0.622 0.862	0.526 0.475 0.798 0.857	NA ~ 0.854 1.007 0.748 0.783	0.748 0.837 NA 1.172 0.971 1.222	1.156 0.984 1.211 < 0.798
<u> </u>	: : 92 F. R. S. P. R. P.		SRH ZUM KGD		: 98 × × × × × × × × × × × × × × × × × ×	VJN XBL :: OL5 P03 STA	01.7 878 U13
Test	09/01/87 05/25/86 02/07/89 03/10/89 12/06/89		11/05/88 12/05/89 10/31/90 09/11/89 05/31/90	07/12/90 09/11/89 04/24/90 07/06/90	02/28/89 02/28/89 03/21/89 04/24/90 07/06/90	04/24/90 VJN 07/06/90 XBL 04/12/88 02/07/89 0L5 03/10/89 P03 12/07/89 STA	02/07/89 03/10/89 12/07/89 02/13/90
Ship- ment	5 5 8 8 5 2	ABEBE 51 51 61	63 33 35 5 64 32 54 55 55 55 55 55 55 55 55 55 55 55 55	67 35 65 65	27 27 27 27 26 65	65 65 65 65 65 65 65 65 65	8 8 8 8 7 8 7 8 7 8 7 8 7 8 7 8 7 8 7 8
No.	2980 2980 2980 2980 2980		3612 3612 3802 3802		3964 3964 3964 4035 4035	4036 4070 4070 4070 4070	7207 7207 7207 7207

Table 9 (Cont'd)

•	() F	* * * * *	* * *	• • •	* * *	* *	+ + +	+ +	* • • •			
	ASSAY	FFFFF	FFF	FER	FFF	FF	***	H		SHEEF	CPE MTT MTT MTT	
	TAI	2.43 1.48 0.00 30.37 6.96	0.00 6.58 14.58	17.50 14.49 24.56	16.81 20.15 12.15	8.97	20.84 25.99 19.77	7.92	10.76 0.62 0.30	1.90° 10.69 13.11 25.48 22.49 10.46	2.40 16.34 11.63 29.08 26.81 14.08	
	S	0.00 0.	0.00	2.29 × 2.83 × 5.08 ×	4.40 × 2.00 × 1.96	1.52 >	5.20 5.45 3.51	1.8. 0.00	2000 2000 2000 2000 2000 2000 2000 200	2.30 2.51 5.29 3.73 0.00	10.00 2.39 2.49 7.53 > 5.55 >	
	A1 95	0.00	0000	0.00	0000	0.00	0.0 10.75 0.00	0.00	9888	0.00000	00.00	
	70 95	0.32 0.32 5.75 1.00 v	1.00	10.00 83.00 88.00	32.00 100.00 100.00	10.00	10.00 32.00 × 32.00	320.00 320.00	%0.60 100.00 100.00	30.30 30.30 10.00 10.00	32.00 - 100.00 100.00 100.00 73.80	
	10 95	0.00 0.00 0.00 0.57 0.00	0.00	0.00	0000	0.00	0.00 > 0.00 > 0.00 >	0.00	00000	0.00 0.00 0.00 0.00 0.00 0.00	32.00 0.00 0.00 0.00 0.00 0.00	
	AI 50	0.00 0.00 5.96 1.43	0.00	2.29 3.98 9.33	10.44	1.52 2.97	7.98 10.11 5.08	3.84	2.67 0.00 0.00 7.36	2.36 3.82 8.20 6.25 0.00	31.00 ~ 6.49 5.64 11.92 8.25 3.18	
	10 50	0.32 0.32 0.74 1.00 v	1.00	10.00 × 23.80 × 20.40	32.00 × 53.70 21.70	10.00 > 20.30	8.89 13.30 8.17	301.00	20.70 100.00 100.00 26.90	3.20 6.60 3.20 2.77 8.30 7.88	100.00 77.10 54.70 83.90 92.70	
	10 50	0.00 0.00 0.17 0.22 v	0.00 0.00 0.22	4.37 > 5.97 2.19	6.41 × 5.14 6.40	6.56 >	1.11	78.50	7.76 0.00 0.00 3.66	2.79 0.84 1.33 0.00	3.20 ~ 17.20 9.69 7.04 11.20 3.02	
	AI 25	1.39 1.05 0.07 8.93 2.43	1.00	6.03 4.14 9.55	6.86 7.42 3.65	2.83 5.08	9.75 10.75 7.30	3.36	4.01 0.00 10.91	10.00 4.27 6.11 12.87 9.07 7.35	32.00 3.94 5.49 13.02 11.34 7.87	
	TC 25	0.32 × 0.03 × 1.00 × 0.32 × 0.	1.00 .	16.90 •	28.20 25.70 12.60	10.00 >	5.80 7.15 5.64	144.00	8.26 0.94 100.00 18.50	3.20 2.80 2.10 1.79 5.03 5.10	32.00 ~ 41.00 24.10 53.00 62.40 6.27	
	10 25	0.23 > 0.31 > 0.47 < 0.11 > 0.13 >	1.00 < 0.67 > 0.09	1.66 v 4.08 1.16	3.46	3.53 >	3.59 0.67 0.77	42.80	2.06 0.00 1.69	0.55	7. 1.00 10.40 4.39 4.07 5.50 0.80	
	Diff.	1.184 0.961 1.252 0.624 0.945	1.039 0.743 0.976	0.748 0.851 0.893	0.742 0.782 0.956	0.742	0.715 0.782 0.899	1.012	0.945 0.818 0.944 1.030	0.852 0.672 0.795 0.818 1.030	0.883 0.672 0.795 0.725 0.977	
	**	/89 0L8 /89 P05 /89 Q68 /89 RJ7 /90 ZXF	/90 TEQ /90 TSP /90 UOX	/89 RL6 /90 VJT /90 XBM	/89 RL7 /89 SPJ /90 ZXG	/89 RL7 /90 ZXG	/89 RL8 /89 SPJ /90 ZXH	/89 ONP /89 R6L	/89 OIN /90 TSQ /90 U14 /90 U0Y	/89 CUC /89 CUC /89 RLA /80 SPK /90 159	/88 /89 0U0 /89 RLA /89 SPK /90 TSR /90 U02	
	Oate	02/07/89 03/10/89 05/23/89 09/11/89 11/01/90	01/18/90 02/06/90 02/13/90	09/12/89 04/24/90 07/06/90	09/12/89 12/06/89 11/01/90	09/12/89 11/01/90	09/12/89 12/06/89 11/01/90	07/06/89 08/15/89	01/31/89 02/06/90 02/13/90 02/13/90	07/27/88 02/27/89 09/12/89 12/06/89 02/06/90	07/28/88 02/27/89 09/12/89 12/06/89 02/06/90	
Shin-	ment	84 84 86 8 84 84 86 86 86 86 86 86 86 86 86 86 86 86 86	62 62	45 65 65	42 61 61	75	42 61 61	33	47 47 63	42 48 48 63 63	48 48 63 63	
AVS	9	4113 4113 4113 4113	4223 4223	4277	4278 4278 4278	4280	4281 4281 4281	4452	4527 4527 4527 4527	0657 0657 0657 0657 0657	4592 4592 4592 4592 4592 4592	

Table 9 (Cont'd)

Confirmatory Assays for Compounds Active Against Yellow Fever Virus

	∢ ∪⊢	• • •	* *	* * • •	• •	+ +	* * *	• • •	* * * *	. +	+ +	• • •	+ + +	
	ASSAY		FF		EE	THE	FFF			ĒĒ	FF			
	¥	12.34 21.70 24.39	6.52	8.03 1.92 0.00 3.54	0.00	5.40	5.05 7.52 3.20	8.37 27.40 31.89	34.86 55.28 45.81 39.69	0.00	9.79	0.55 0.00 14.42	4.01 0.98 3.94	
	IS	1.71 * 4.16 * 4.23	0.00 >	0.00	0.00	1.34	0.00	8.0% 9.0% \$.0%	4.04 > 23.82 > 21.59 > 8.37 >	0.00	1.79 >	2.00	0.38	
	AI 95	0.00	0.00	00000	0.00	0.00	0.00	0.00	3.57 10.91 > 0.00 > 3.45 >	0.00	1.06 > 3.20	0.00	0.00	
	26 21	0.10	320.00	247.00 100.00 99.10 95.30	9.86	305.00	320.00 32.00 32.00	320.00 100.00 320.00	320.00 > 320.00 > 320.00 > 100.00 >	320.00	320.00 > 966.00	1000.00 1000.00 3200.00	320.00 983.00 320.00	
	S6 21	0.00	0.00	00000	0.00	0.00	0.00	0.00.0	89.60 × 29.30 × 0.00 × 29.00 ×	0.00	302.00 >	0.00	0.00	
•	AI 50	1.71 6.97 8.31	0.00	0.00	0.00	2.31	0.00	3.67 16.45 14.89	8.58 23.82 21.59 8.37	0.00	3.69	0.00	1.39	
	10 50	0.10 > 0.31	320.00	67.60 62.00 57.70 52.80	6.71	173.00 320.00	16.60 18.50 8.09	79.60 100.00 > 320.00 >	171.00 320.00 > 320.00 > 100.00 >	320.00 806.00	320.00 >	958.00 564.00 2890.00	320.00 > 253.00 238.00	
	10 50	0.06 0.03 v	0.00 >	29.00 0.00 0.00	0.00	74.90	0.00 0.00 3.20	21.70 6.08 > 21.50 >	20.00 13.40 > 14.80 >	0.00 >	179.00 >	0.00	230.00 > 0.00 0.00	
	AI 25	2.57 9.21 11.96	4.53	3.06 1.86 0.00 0.92	0.00	2.04	3.77	3.24 11.18 14.14	8.07 44.31 40.85 17.38	9.20	2.39	0.00	1.13 2.06 0.00	
	10 25	0.10 > 0.18 0.13	320.00 > 712.00	47.80 39.90 34.80 29.40	4.95	100.00	6.60 7.00 2.83	21.50 49.00 207.00	80.80 320.00 > 320.00 > 100.00 >	320.00 459.00	320.00 > 490.00	426.00 215.00 1370.0	87.10 66.00 130.00	
	10 25	0.02	70.70 >	15.60 21.50 0.00 32.00	5.01	00.0	3.20 1.86 1.65	6.63 4.38 14.70	10.00 7.22 > 7.83 > 5.76 >	40.00 4	134.00 > 134.00	0.00	77.20 32.00 0.00	
	biff.	0.892 0.765 0.944	0.822	0.749 1.023 0.791 0.966	0.621	0.816	0.993 0.577 0.899	0.943 0.732 0.893	1.222 0.751 0.765 0.986	0.783	0.765	0.725 1.220 0.848	0.919	
	± *	SRI U10	VLV	RAL UF9	100 100	02K U16	OFN RLC ZXH	OFS RLD SPL	RLG SPO 1SA	P13 SRJ	P1A SP0	SRC TO2 TVR	CUB RU9 S23	
	Test	02/27/89 12/05/89 02/13/90	04/24/90	07/06/89 08/15/89 03/08/90 05/03/90	11/02/88	11/08/88 02/13/90	01/24/89 09/12/89 11/01/90	01/24/89 09/12/89 12/06/89	02/21/89 09/12/89 12/06/89 12/06/90	03/13/89 12/05/89	03/13/89 12/06/89	12/05/89 12/20/89 02/08/90	02/27/89 10/02/89 10/10/89	
	Ship- ment	8 8 5	59 59	3333	77	33	999	97 97 97 97 97 97 97 97 97 97 97 97 97 9	48 48 48 48	27	15	62 45 GABSN	8 8 8 8 7 8 7	
	AVS.	6097	4611	4736 4736 4736 4736	137	4768	4785 4785 4785	9627	4855 4855 4855 4855	8267	7665	5040 5040 5040	5072 5072 5072	

Table 9 (Cont'd)

∢ ∪⊢	+ + +	+ + +	+ + +	+	+ +	+ + +	* * *	* * *		+ 1 +	
ASSAY	# # #	H H H	HAM	THE REPORT OF THE PERSON OF TH	FE	E E E	M M M	FFF	MANA	FF F	
TAI	25.50 20.19 37.43	18.57 27.36 24.85	0.37 10.12 0.56	0.00 0.38 6.76	13.17	0.00 0.00 17.25	4.37 30.16 24.30	5.94 13.16 16.73	6.19 8.33 11.88 26.75	7.36	1.11 4.39 11.55 5.82
S	4.96 3.22 × 8.30 ×	2.59 × 5.90 4.61 ×	0.00 2.63 0.00	0.00	3.63	0.00	0.00 11.96 v	3.36	0.00 0.93 0.00 0.00 0.00 0.00 0.00 0.00	1.73 > 0.00 > 1.49 >	0.00
A1 %	3.47	1.10 > 3.5 1.21 >	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.000
70 %	309.00 309.00 320.00 >	320.00 > 1000.00 > 1000.00 >	320.00 1000.00 1000.00	274.00 296.00 307.00	320.00 966.00	10.00 6.91 8.19	320.00 100.00 100.00	320.00 83.00 9.24	320.00 1000.00 1000.00 320.00	320.00 1000.00 320.00	1000.00 100.00 32.00
10.95	89.00 0.00 97.90 v	291.00 > 283.00 > 827.00 >	0.00	0.00	0.00	0.00	0.00	0.00	0.0000	0.00	0.0000
AI 50	6.72	2.59 8.45 4.61	0.00 0.00 0.00	0.00	0.00	5.00	0.00 24.42 5.85	0.00 5.10 5.34	0.00 1.02 0.00 4.17	1.73	0.00 2.68 0.00
10 50	210.00 210.00 320.00 >	320.00 * 806.00 * 1000.00 *	312.00 341.00 311.00	71.90 89.90 159.00	320.00 578.00	1.00	20.40 65.30 29.00	5.47 2.69 0.24	320.00 1000.00 > 698.00 320.00 >	320.00 > 711.00 320.00 >	682.00 6.78 6.08 2.34
10 50	31.20 48.10 38.60 >	123.00 > 95.40 217.00 >	0.00	0.00 0.00 86.70	0.00 100.00	0.00	0.00 2.68 4.95	0.00	0.00 y 976.00 y 0.00 y 76.70 y	185.00 > 0.00 > 215.00 >	0.00 2.27 0.00
VI 25	8.98 5.08 15.74	4.98 10.68 6.79	0.56 0.00 0.00	0.00	6.76	1.00	0.96 25.21 14.57	2.42 5.53 5.20	0.00 1.70 6.39 11.81	2.68 0.00 2.18	0.00 2.84 2.74 2.74
1C 25	155.00 155.00 320.00 >	. 320.00 > 563.00 . 1000.0 >	100.00 197.00 194.00	45.00 30.20 85.10	> 320.00 > 363.00	* 1.00 * 0.45	4.23 32.00 19.50	2.42 > 1.77 > 0.17 >	320.00 906.00 509.00 320.00 >	> 320.00 > 478.00 > 320.00 >	3.80 2.96 1.35
10 25	17.30 30.50 20.30 >	64.20 > 52.70 147.00 >	179.00 49.00 0.00	0.00	47.30 3	9.00	4.39	1.00	0.00 532.00 79.60 27.10 	0.00	0.00 1.34 0.65 0.49
Diff.	0.935 0.892 0.856	1.012 0.940 0.777	0.952 0.877 0.776	1.326 0.920 0.845	0.800	1.071 0.832 1.155	1.235 0.708 0.871	1.059 < 0.781 < 0.866 <	1.109 0.912 0.904 0.781	1.142	1.119
i *	R18 2X1	A M N	V S N N	RUB 284	SXA	PC8 VY0 X9E	VNP XBO	PCA VYO XJX	SYK SXS VY0		R3E 2XJ
Test	06/20/89 08/07/89 11/01/90	07/10/89 08/15/89 04/26/90	07/24/89 10/03/89 04/26/90	07/24/89 10/03/89 10/04/90	03/07/89	03/27/89 05/08/90 07/06/90	04/11/89 04/26/90 07/06/90	03/27/89 05/08/90 07/18/90	03/27/89 12/12/89 12/14/89 05/08/90	06/13/89 08/07/89 06/13/89	08/07/89 06/13/89 08/07/89 11/01/90
Ship-	222	57 57 65	8 8 8	8 8 8	25	223	65 53	288	8888	2 2 2	2 222
AVS	5121 5121 5121	5138 5138 5138	5186 5186 5186	5197 5197 5197	5241	5405 5405 5405	5450 5450 5450	5483 5483 5483	7875 7875 2787 2787	5535 5535 5538	5538 5539 5539 5539

Table 9 (Cont'd)

Confirmatory Assays for Compounds Active Against Yellow Fever Virus

	∢ ∪⊢	+ + + +		+ +	• • • •	+ +	+ +	* * *	* * *	* * *	+ +	+ •	+ +
	ASSAY		EEEE			E E	EE		FFF	EEE		EE	T I
	TAI	1.85 5.43 0.00 7.77	13.40 5.77 0.00	3.01	3.03 0.00 14.50		17.81 21.58	11.86 30.66 21.00	0.21 8.63 17.53	23.80 26.85 20.89	19.66	3.49	32.24
	ïs	0.00 1.03 1.36	0.00		0.94 v 0.00 v 0.03 v	1.08 •	3.18 >	3.12 > 8.34 > 4.56	0.00 4.90 0.00	4.33 × 5.89 3.47	3.06 ×	0.00	6.15 > 2.13 >
	A1 95	0.000	0.000	0.00	80 00	0.0	0.00	0.00 0.00 0.00	0.00	1.20 × 0.00 3.28	3.19	0.00	3.29 > 1.08 >
	26 21	320.00 32.00 32.00 100.00	1000.00 3200.00 1000.00 3200.00	320.00	9.38 9.38 94.70 91.90	91.80	320.00 100.00	320.00 32.00 32.00	9.13 0.97 3.15	320.00 > 964.00 966.00	320.00 > 956.00	320.00 313.00	320.00 >
	10 95	9888	9888	A A	86 86	0.00	0.00	0.00	9000	267.00 > 0.00 295.00	286.00 > 300.00	0.00	97.30 > 297.00 >
,	A1 50	0.00 2.12 0.00 2.77	0.00	1.93	5.30	2.71	11.82	9.73 15.92 10.15	0.00	4.33 4.68	3.06	1.31	6.15
	TC 50	17.10 1.59 32.00 77.90	873.00 3200.00 > 916.00 1670.00	320.00 >	16.10 3.82 46.90	22.20	52.40 32.00	9.73 > 32.00 > 27.70	2.53	320.00 > 638.00 660.00	320.00 > 564.00	320.00 > 212.00	320.00 >
	10 50	0.00 0.75 0.00 v	3720.00 × 0.00 0.00 0.00	144.00 > 254.00	8.85 0.00 8.85 0.00	8.19	4.43	2.01 2	0.00	74.00 v 77.60 141.00	105.00 > 165.00	245.00 > 0.00	52.00 > 150.00 >
	AI 25	1.44 1.88 0.00 2.60	0.00 0.00 0.90	1.68	1.56 1.13 7.56 7.56	2.21	14.07	3.12 < 18.09 16.03	1.00	6.58 9.98 5.73	5.52	20.0	8.70 3.10
	10 25	8.91 0.77 32.00 38.30	95.80 2710.0 618.00 606.00	2 2	8.30 2.22 24.20 19.50	8.87	14.10 >	3.12 > 16.80 12.40	1.00 > 0.49 > 1.57	320.00 > 457.00 490.00	320.00 > 346.00	220.00	320.00 >
	10 25	6.18 0.41 14.70	0.00 280.00 0.00 674.00	68.40 > 123.00	3.20	4.02	1.00	< 1.00 0.93 0.78	1.000.100.13	48.60 × 45.80 85.50	57.90 × 119.00	134.00	36.80 >
	Diff.	1.042 1.101 0.968 0.810	0.821 0.897 0.915 0.853	0.989	0.738	0.795	0.712	0.712 0.539 0.886	0.984 0.857 0.732	0.703 1.087 0.911	0.703	0.712	0.712
	¥ *	00E R3G VSN 28X	SRE TVT UGF ZXK		SSE 358		SEF ZXL	SEF 101 1VK	SVA TVL WPE	70C 2UI 1SB	10C	10E	158
	Test	06/13/89 08/07/89 05/01/90 10/04/90	12/05/89 02/08/90 05/31/90 11/01/90	07/17/89	10/30/89 01/30/90 10/23/89 01/23/90	11/07/89	11/07/89	11/07/89 02/01/90 02/08/90	12/12/89 02/08/90 06/13/90	02/01/90 10/31/90 12/06/90	02/01/90 10/31/90	02/01/90 10/31/90	02/01/90 TOE 12/06/90 1SB
	Ship- ment	2252	62 GABSN 67 62/67	57 57	99 99	22	2 5	19 19	19 19	2 2 2	23	29	62
	AVS No.	5543 5543 5543 5543	5601 5601 5601 5601	1695	5905 5905 5916 5916	5995 5995	5997 5997	5998 5998 5998	7709 7709 7709	6195 6195 6195	6196 6196	6199	6200

Table 9 (Cont'd)

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ASSAY	FF	T TH	FF	FF	FF	ĒĒ	ĒĒ	ĔĔ	FF	FF	FFFF	E E	ĒĒ	ĒĒ	H
TAI	32.24 20.26	10.98 22.15	19.96 15.14	0.0	10.18	24.36	43.56	24.45	26.49	47.17	0.26 4.57 0.00 20.74	17.65	16.55	0.92	2.19
15	6.07 > 3.27 >	3.17	2.74	0.55	1.81 > 2.74	4.33 >	17.12 > 17.09 >	4.41 > 5.83 >	6.16 >	18.33 >	0.00	2.18 > 0.00	3.51	0.00	0.00
A1 95	3.41 > 1.13 >	0.00	3.27	0.00	1.06 >	1.20 >	7.22 > 3.20 >	1.20 >	3.42 > 0.00	10.62 >	0.000	0.00	0.00	0.00	0.00
8 21	320.00 >	320.00 966.00	309.00	320.00 88.20	320.00 > 966.00	320.00 >	320.00 >	320.00 >	320.00 >	320.00 > 966.00	1000.00 3200.00 3200.00 3200.00	320.00 320.00	95.10	320.00 925.00	30.90
96 JI	93.80 > 284.00 >	0.00	94.50	9.00	302.00 >	267.00 > 94.50 >	44.30 >	267.00 > 94.20 >	93.70 >	30.10 >	00000	0.00	0.00	0.00	0.0
AI 50	3.27	4.28	3.71	0.76	1.81	4.33	17.12	5.83	6.16	3.94	0.00	3.24	6.85	0.00	0.00
10 50	320.00 >	320.00 >	210.00	22.80	320.00 >	320.00 > 694.00	320.00 >	320.00 >	320.00 > 927.00	320.00 × 660.00	1000.00 3200.00 3040.00 3200.00 >	286.00 282.00	50.70	294.00	21.10
10 50	52.70 > 97.80 >	179.00 >	56.60	30.00	177.00 \$	74.00 >	18.70 >	72.60 > 54.90 >	\$2.00 > 74.90	17.50 >	0.00 v 0.00 v 0.00 v 1530.00 v	88.50 0.00	7.41	0.00	0.00
AI 25	8.36	2.49	3.6	0.00	3.66	6.58	23.41	6.88	8.55	24.81	0.00 2.24 0.63 5.05	6.04	7.43	0.00	3.05
10.25	320.00 >	. 320.00 >	155.00	16.40	. 320.00 >	320.00 > 507.00	320.00 >	320.00 >	320.00 > 624.00	. 320.00 >	1000.0 3200.0 > 2020.0 3200.0 >	193.00	26.00	92.70 35.70	15.60
S 21	38.30 >	129.00 >	42.50	17.30	131.00 >	48.60 >	13.70 >	46.50 >	37.40 >	12.90 >	0.00 > 1430.0 > 3200.0 634.00 >	32.00	3.50	0.00	0.00
Diff.	1.009	0.767	0.740	0.740	0.720	1.020	0.755	0.994	0.994	0.996	0.613 0.928 0.827 0.553	0.669	0.598	0.840	0.611
<u>=</u> *	10F 1SC	10F 2U.S	10G	70G ZUK	101 20L	TBR ZXL	181	18U	18 1 3 5 1	TBX ZXM	SRF TVU WGG ZXN	SI B	SIE	200	X 700
Test	02/01/90 TOF 12/06/90 1SC	02/01/90	02/01/90	02/01/90	02/01/90	01/16/90	01/16/90	01/16/90	01/16/90	01/16/90	12/05/89 02/08/90 05/31/90 11/01/90	11/14/89 02/06/90	11/14/89 02/06/90	02/15/90 03/20/90	02/15/90
Ship- ment	33	33	33	33	22	33	33	33	22	22	62 GABSN 67 67	4 4	5 5	22	28
AVS.	6201	6202	6203	6204	6207	6214 6214	6218 6218	6219	6220	6225	6234 6234 6234 6234	6250	6256	6309	6315

Table 9 (Cont'd)

Confirmatory Assays for Compounds Active Against Yellow Fever Virus

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ASSAY	ĒĒ	ĒĒ	EEE	E E	ĒĒ	ĒĒ	ĒĒ	ĒĒ	EE	ĒĒ	ĒĒ	ĒĒ	ĒĒ	EE	FE
¥	11.77	3.67	3.78 21.91 0.00	7.19	6.71	7.30	19.99	2.92	12.50 28.19	2.50	4.52	19.18	0.00	36.65	6.53
15	0.00	1.75 >	3.47 >	1.84	0.00	0.00 > 13.77 >	2.53 > 3.27 >	0.00	0.00 >	3.54 >	3.00	5.01 >	1.60 \$	6.79	0.00 >
¥ %	0.0	0.00	0.00 3.28 0.00	0.00	0.00	0.00	0.0	0.00	0.00	9.38	0.0	0.00	0.00	0.00	0.0
70 %	163.00	303.00	96.40 96.60 80.00	320.00	320.00	320.00	96.50 95.30	210.00	320.00	320.00	99.00	1000.00	1000.00	3200.00	100.00
5 %	0.00	0.00	0.00 29.50 0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00 >	0.00	0.00	0.00	0.00	0.00
A1 50	0.00	1.58	0.00	0.00	0.00	0.00	3.45	3.25	0.00	5.07	2.23	5.01	9.0	0.00	0.00
10 50	67.20	148.00	2.38 5.00 7.00 7.00	320.00	217.00 657.00	320.00 731.00	65.30 52.80	63.20	320.00	266.00 251.00	54.70	1000.00 > 3200.00	1000.00 >	1000.00 >	67.80 95.00
10 50	8.0	93.90 3.09	0.00	0.00 >	313.00	29.80	18.90 9.06	61.00	0.00 >	0.00	23.80	200.00 >	626.00 >	147.00 > 0.00 >	12.10
A1 25	5.68	2.24	2.36 5.80 0.00	3.49	0.00	17.89	21.56 12.63	1.95	3.99	5.04	1.92	12.09	2.24	123.02	5.66
10 25	10.40	50.10	46.30	272.00	0.96	320.00 >	48.00	40.30	320.00 > 1000.0 >	181.00 175.00	29.70	1000.0 > 2510.0	1000.0 > 3200.0	1000.0 > 3200.0	63.50
10 23	8.39	22.40	19.70 8.45 0.00	77.90	0.00	17.90 >	2.22	20.70	80.20 > 36.10 >	34.80	15.40	82.70 >	< 00.0	8.13 > 0.00 >	8.81 5.56
Diff.	0.809	0.795	0.896 0.871 0.553	0.741	0.890	0.677	0.979	1.021	0.945	0.900	0.804	0.760	0.574	0.574	0.785
ž*	0 US	0 UKP	0 U61	O VIE	0 M36	0 UAR	O UAR	O UCT	DO CC	200	0 K37	0 VAI	0 VF5	0 VF5	NO WGH
Test	02/20/90 U4L 03/20/90 U02	02/20/90	02/27/90 03/20/90 11/01/90	05/10/90	05/15/90 436 07/06/90 xBV	03/01/90 UAR 03/22/90 UQ9	03/01/90	03/06/90 UCT 04/18/90 VAT	03/06/90 UCU 03/22/90 UGC	03/06/90 UCV 03/22/90 UGD	05/15/90 07/11/90	04/18/90 VAI 06/27/90 X3T	04/19/90	04/19/90 VFS 06/27/90 X41	05/31/90 WGH 07/12/90 XFU
Ship- ment	22	22	222	33	33	22	23	22	63	23 23	88	22	22	22	19
AVS.	6326	6334	6369 6369 6369	222	6422 6422	***	6445	6456	6458 6458	33	£ 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2	6589	6617 6617	8139 8139	6707 6707

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Table 9 (Cont'd)

Confirmatory Assays for Compounds Active Against Yellow Fever Virus

* *	+ +	+ +	+ +	* *	+ +	+ +	• •	+ •	+ +	+ •	• •	• •	* * *	* * *	
H	EE	EE	FF	FF	MTT	EE	FIE	EE	ĒĒ	EE	ĒĒ	ĒĒ	FFF	EEE	
47.19	9.56	3.02	30.64	9.53	17.36	12.38	3.73	6.98	13.35	1.67	9.27	1.02	0.11 7.19 13.92	9.34 33.97 32.54	
19.75 × 9.11 ×	2.01 > 3.13 >	3.12 >	5.83 > 6.12	2.22 5.73 ×	3.62	2.74	0.00	1.39 >	1.54 > 0.00 >	0.37 >	0.00	0.00 >	0.10 1.84 2.20 v	0.26 9.86 8.77 ×	
35.49	0.00	0.00	3.40 > 0.00	3.47	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	31.97	
320.00 >	24.10 32.00	320.00	320.00 > 955.00	320.00	320.00 965.00	320.00	320.00	320.00	310.00	320.00	320.00	320.00	320.00 918.00 869.00	320.00 906.00 320.00	
9.02 \$	0.00	715.00 >	94.20 >	0.00 > 279.00	0.00	9.00	0.00	0.00	0.00	0.00	0.00 > 945.00 >	0.00	0.00	0.00 28.30 0.00 ×	
69.50	2.94	0.00	5.83 10.28	3.13	3.62	4.02	0.00 2.21	1.39	2.31	0.00	0.00	0.00	1.30	7.14 22.95 31.33	
247.00	6.86	320.00	320.00 > 553.00	238.00	320.00 > 653.00	255.00	320.00	320.00 > 667.00	193.00	320.00 > 163.00	320.00	320.00	130.00 94.00 81.80	155.00 218.00 192.00	
3.55	2.33	0.00 > 186.00	54.90 >	75.90 85.50	88.50 > 0.00	63.60	0.00 > 252.00	230.00 > 0.00	63.70	320.00 > 0.00	0.00 > 566.00 >	0.00 > 759.00 >	100.00 23.90 18.30	21.70 9.48 6.12	
55.53	4.68 5.59	1.21	7.87	3.42	10.00	3.87	2.07	2.11	3.74	1.29	0.00	0.00	3.22	0.49 19.13 15.33	
70.10	4.66 4	320.00 > 582.00	320.00 > 329.00	169.00	320.00 >	174.00	320.00	320.00 > 493.00	128.00 75.60	117.00	320.00	320.00	10.30	5.75 93.50 53.60	
1.28	1.00	264.00 >	40.70 > 37.50	49.30	32.00 > 294.00	45.10	0.00 >	151.00 >	34.40	90.90	0.00 >	493.00	41.10 13.70 9.45	11.80 4.89 3.50	
0.877		0.990	0.990	1.062	0.729	0.750	1.362	1.119	0.984	1.000	1.030	0.743	1.463 1.164 1.247	1.444	
	WPO XG3	XO4	X X Z	XSV	X19	X10 X2E	XSX	XV4 YMS	XYY	XY2 YPX	XY4 YPZ	YCC 26N	YE9 26P 2P0	7EA	
06/05/90	06/13/90	07/25/90	06/90/60	07/26/90	06/21/90	06/21/90	07/26/90	06/01/90	08/02/90	08/02/90	08/02/90	08/30/90	08/31/90 10/03/90 10/25/90	08/31/90 10/03/90 10/25/90	
79	79	55	99	69		33	\$\$	5 5	69	69	69	22	222		
7219	6792	6942	6943	9769	6269	9869	7003	7032	7044	7045	7049	7051	7065 7065 7065	7067 7067 7067	
	67 06/05/90 WIY 0.877 1.26 70.10 55.53 3.55 247.00 69.50 9.02 > 320.00 > 35.49 19.75 > 47.19 67 07/12/90 XFW 0.657 1.28 17.90 14.05 1.97 > 320.00 > 162.60 0.00 > 320.00 0.00 9.11 > 26.68	67 06/05/90 WIY 0.877 1.26 70.10 55.53 3.55 247.00 69.50 9.02 > 320.00 > 35.49 19.75 > 47.19 67 07/12/90 XFW 0.657 1.28 17.90 14.05 1.97 > 320.00 > 162.60 0.00 > 320.00 0.00 9.11 > 26.68 67 06/13/90 WPD 0.894 < 1.00 4.68 > 4.68 2.33 6.86 2.94 0.00 24.10 0.00 2.01 > 9.56 67 07/12/90 XG3 0.675 1.23 6.90 5.59 2.21 13.00 5.91 0.00 > 32.00 0.00 3.13 > 17.23	67 06/05/90 WIY 0.877 1.26 70.10 55.53 3.55 247.00 69.50 9.02 > 320.00 > 35.49 19.75 > 47.19 67 07/12/90 xFW 0.657 1.28 17.90 14.05 1.97 > 320.00 162.60 0.00 > 320.00 0.00 9.11 > 26.68 67 06/13/90 WPO 0.894 < 1.00 4.68 > 4.68 2.33 6.86 2.94 0.00 24.10 0.00 2.01 > 9.56 67 07/12/90 xG3 0.675 1.23 6.90 5.59 2.21 13.00 5.91 0.00 > 32.00 0.00 3.13 > 17.23 69 07/25/90 xQ4 0.990 264.00 > 320.00 > 1.21 0.00 > 320.00 0.00 > 320.00 0.00 3.13 > 17.23 69 09/06/90 YMI 0.970 124.00 582.00 4.70 186.00 843.00 4.52 715.00 > 1000.00 > 1.40 3.12 > 22.51	67 06/05/90 WIY 0.877 1.26 70.10 55.53 3.55 247.00 69.50 9.02 > 320.00 > 35.49 19.75 > 47.19 67 12/90 XFW 0.657 1.28 17.90 14.05 1.97 > 320.00 162.60 0.00 > 320.00 9.11 > 26.68 67 07/12/90 XFW 0.657 1.28 17.90 14.05 1.97 > 320.00 162.60 0.00 24.10 0.00 9.11 > 26.68 67 07/12/90 XGJ 0.675 1.23 6.90 5.59 2.21 13.00 5.91 0.00 320.00 0.00 3.13 > 17.23 6.90 07/12/90 XGJ 0.90 264.00 > 320.00 1.21 0.00 > 320.00 0.00 0.00 3.10 3.12 > 22.51 69 09/06/90 YMI 0.970 124.00 582.00 4.70 186.00 843.00 4.52 715.00 > 1000.00 1.40 3.12 > 22.51 69 09/06/90 YMI 0.986 37.50 329.00 8.77 53.80 533.00 10.28 0.00 955.00 0.00 55.00 56.20 27.23	67 06/05/90 WIY 0.877 1.26 70.10 55.53 3.55 247.00 69.50 9.02 > 320.00 > 35.49 19.75 > 47.19 67.19 67.12/90 xfw 0.657 1.26 17.90 14.05 1.97 > 320.00 > 162.60 0.00 > 320.00 9.11 > 26.68 67 07/12/90 xfw 0.657 1.25 17.90 14.05 1.97 > 320.00 > 162.60 0.00 > 320.00 9.11 > 26.68 67 07/12/90 xd3 0.675 1.23 6.90 5.59 2.21 13.00 5.91 0.00 24.10 0.00 2.01 > 9.56 67 17.25 0.00 24.10 0.00 2.01 0.00	67 06/13/90 kHY 0.877 1.26 70.10 55.53 3.55 247.00 69.50 9.02 > 320.00 > 35.49 19.75 > 47.19 26.68 67 107/12/90 xHy 0.657 1.28 17.90 14.05 14.05 1.97 > 320.00 > 162.60 0.00 > 320.00 > 320.00 9.11 > 26.68 67 07/12/90 xG3 0.675 1.23 6.90 5.59 2.21 13.00 5.91 0.00 24.10 0.00 24.10 0.00 2.01 > 9.56 07/12/90 xG3 0.675 1.23 6.90 5.59 2.21 13.00 5.91 0.00 320.00 0.00 0.00 37.13 > 17.23 69/06/90 xHI 0.970 124.00 582.00 4.70 128.00 5.20 1.21 0.00 520.00 0.00 0.00 520.00 520.00 0.00	67 06/12/90 WHY 0.877 1.28 70.10 55.53 3.55 247.00 69.50 9.02 > 320.00 > 35.49 19.75 > 47.19 67 06/13/90 WHO 0.894 < 1.00 4.68 > 4.68 2.31 6.86 2.94 0.00 24.10 0.00 2.01 > 9.11 > 26.68 67 07/12/90 XF4 0.890 2.64.00 > 320.00 1.21 13.00 > 320.00 0.00 24.10 0.00 2.01 > 9.56 69 07/25/90 XG4 0.990 2.64.00 > 320.00 1.21 0.00 > 320.00 0.00 0.00 2.01 > 9.50 69 07/25/90 XG4 0.990 40.70 320.00 7.87 320.00 320.00 3.40 3.40 3.12 > 22.51 69 07/25/90 XG4 0.990 40.70 320.00 7.87 32.80 320.00 3.40 320.00 3.40 3.12 > 22.51 69 07/25/90 XG4 0.990 40.70 320.00 7.87 32.80 320.00 3.13 0.00 320.00 3.40 3.12 > 22.51 69 07/25/90 XG4 0.990 40.70 320.00 7.87 32.80 320.00 3.13 0.00 320.00 3.40 3.12 > 22.51 69 07/25/90 XG4 0.990 40.70 320.00 3.42 32.80 330.00 3.13 0.00 320.00 3.40 3.12 > 22.51 69 07/25/90 XG4 0.990 40.70 320.00 3.42 32.80 330.00 3.42 320.00 3.40 3.12 > 22.51 69 07/25/90 XG4 0.990 40.70 320.00 3.42 32.80 3.43 3.40 3.40 3.40 3.40 3.12 > 22.51 69 07/25/90 XG4 0.990 40.70 320.00 3.42 320.00 3.40	67 06/13/90 WIY 0.877 1.28 70.10 55.53 3.55 247.00 69.50 9.02 > 320.00 > 35.49 19.75 > 47.19 67 06/13/90 WIY 0.657 1.28 17.90 14.05 14.05 1.97 > 320.00 > 162.60 0.00 > 320.00 > 35.49 19.75 > 47.19 67 06/13/90 WIY 0.657 1.28 17.90 14.05 2.33 6.86 2.34 0.00 24.10 0.00 20.00 9.11 > 26.68 67 06/13/90 WIY 0.894 1.02 6.90 5.50 1.21 0.00 > 320.00 1.00 0.00 24.10 0.00 24.10 0.00 3.13 > 17.23 68 06/21/90 WIY 0.976 124.00 582.00 1.23 1.65.00 532.00 1.00 0.00 0.00 3.40 0.00 0.00 3.13 > 17.23 69 07/25/90 WIX 0.986 2.23 490.00 7.87 54.90 532.00 5.83 94.20 \$320.00 3.40 5.53 30.44 69 07/25/90 WIX 0.986 52.30 490.00 9.37 55.90 532.00 \$3.13 0.00 \$320.00 3.40 5.53 \$30.44 69 06/21/90 WIX 0.988 52.30 490.00 9.37 85.90 532.00 \$3.13 0.00 \$320.00 3.47 5.73 \$7.01 68 06/21/90 WIX 0.988 52.30 480.00 1.63 0.00 653.00 0.00 0.00 320.00 0.00 0.00 3.47 5.73 \$7.01 68 06/21/90 WIX 0.988 52.30 480.00 1.63 0.00 653.00 0.00 0.00 320.00 0.00 0.00 0.00 0.0	67 07/12/90 MH 0.877 1.28 70.10 55.53 3.55 247.00 69.50 9.02 > 320.00 > 35.49 19.75 > 47.19 6.65 07 10.00 2.01 > 320.00 46.26 0.00 > 320.00 9.11 > 26.66 6.7 07/12/90 MH 0.86	67 07/12/90 kH 0.657 1.28 17:90 14:05 13.55 247.00 69.50 9.02 320.00 9.00 9.15.68 66 66 67 07/12/90 kH 0.657 1.28 17:90 14:05 17:00 14:05 17:00 14:05 17:00 14:05 17:00 14:00 14:05 17:00 14:00	67 07/12/90 NIA 0.687 1.28 17.90 14.05 14.05 14.05 19.05 182.00 182.00 182.00 182.00 182.00 19.05 19.0	67 06/12/90 KHV 0.687 1.28 177.90 14.05 1.35 247.00 69.50 9.02 320.00 9.55.9 19.75 47.19 26.68 67 07/12/90 KHV 0.657 1.28 177.90 14.05 14.05 1.37 520.00 162.60 0.00 320.00 9.11 26.68 67 07/12/90 KHV 0.985 1.02 1.28 177.90 14.05 1.29 1.30 0.00 32.00 0.00 0.00 32.00 0.00 0.00	64 06/15/90 WIY 0.687 1.26 70.10 55.53 3.55 247.00 162.60 0.00 530.00 55.49 19.75 47.19 26.66 67 07/12/90 XRU 0.657 1.28 17.79 14.05 14.05 14.05 14.05 14.05 0.00 530.00 0.00 0.00 0.01 5.66 0.00 530.00 0.00 0.00 0.01 5.66 0.00 530.00 0.00 0.00 0.01 5.66 0.00 530.00 0.00 0.00 0.01 5.66 0.00 530.00 0.00 0.00 0.01 5.66 0.00 530.00 0.00 0.00 0.00 0.01 5.66 0.00 530.00 0.00 0.00 0.00 0.00 0.00 0.	67 06/12/90 km 0.894 < 1.0.2 17.90 14.05 1.97 > 320.00 162.60 0.00 320.00 35.40 19.75 > 47.19 68 06/12/90 km 0.687 1.0.2 17.90 14.05 1.97 > 320.00 162.60 0.00 320.00 0.00 2.01 > 56.60 69 06/12/90 km 0.695 2.0.0 320.00 1.0.1 1.20 1.97 > 320.00 162.60 0.00 320.00 0.00 2.01 > 56.60 69 06/12/90 km 0.995 2.0.0 320.00 1.21 0.00 320.00 0.00 320.00 0.00 3.10 0.00 3.10 69 06/12/90 km 0.995 2.0.0 320.00 1.22 0.00 320.00 0.00 1.00 1.00 320.00 0.00 1.00 1.00 1.00 69 06/12/90 km 0.995 2.0.0 320.00 1.21 0.00 320.00 1.22 0.00 320.00 0.00 1.00 1.00 1.00 1.00 1.00 1.00	67 00005590 MIN 0.687 1.28 70.10 55.53 1.37 327.00 69.30 0.00 320.00 0.00 50.00 50.10

Table 9 (Cont'd)

Confirmatory Assays for Compounds Active Against Yellow Fever Virus

	∢ ∪⊢	* * *	. + +	* * *	* * *	* * *	• • •			+ •	• • • •		* *
	ASSAY	FFF		FFF			EEE		# # # #	T T			F F
	TAI	8.13 19.91 26.73	0.00 1.34 10.07	17.13 15.83 7.35	13.21 5.56 11.84	19.39 4.83 16.05	8.07 4.42 8.12	24.20 7.65 12.79	0.03 22.36 20.15	18.82	15.50	18.62 2.57	3.35
	5	2.05 3.83 5.62 >	0.00	2.37 > 2.34 0.53 >		3.49	0.00	4.65 0.00 0.00	0.00 × 5.68 × 2.11 ×	4.07	0.00 4	0.74 > 0.85	1.20 >
	VI 95	9.51 0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	88 88	0.00	0.00
	TC 95	306.00 293.00 320.00	100.00 869.00 959.00	309.00 305.00 313.00	100.00 %.40 100.00	100.00 298.00 100.00	100.00	100.00	320.00 1000.00 1000.00 >	32.00	320.00 320.00 320.00	320.00	320.00 938.00
	10 %	30.80	0.00	0.00	0.00	0.00	9000	0.00	0.00 > 0.00 > 984.00 >	0.00	00 00 88 88	0.00	0.00
	AI 50	3.34 6.56 10.29	0.00	6.69 4.13 1.86	3.71	8.92 0.00 2.80	1.56	8.10 0.00 0.00	5.68	7.58	6.34 0.00 2.07	0.74	1.27
	10 50	181.00 87.00 189.00	100.00 216.00 495.00	147.00 171.00 145.00	27.60 41.70 44.60	56.30 59.20 59.20	10.00 24.90 26.20	31.50 56.00 40.20	320.00 1000.00 > 1000.00 >	21.60	320.00 207.00	320.00 >	382.00
	10 50	54.10 13.20 18.40	0.00 > 226.00	21.90 41.50 77.60	0.00	6.31 0.00 21.10	0.00	3.88	0.00 > 176.00 > 473.00 >	2.85	100.00	432.00 > 310.00	252.00 > 0.00
and I have	AI 25	3.80 7.88 9.05	0.00	5.20 7.17 2.04	7.34 2.30 5.03	6.39 2.11 10.95	1.91 2.40 2.98	13.33 3.13 5.26	8.32 3.44	8.42	7.41 0.00 3.22 1.00	10.52	1.90
	TC 25	50.80 50.80 103.00	24.70 109.00 227.00	52.00 96.90 41.10	13.90 21.30 20.20	22.00 23.10 33.00	6.60 16.40 15.80	18.00 21.80 19.30	320.00 1000.0 >		320.00 128.00	1.1.4	302.00 137.00
	10 25	29.30 6.45 11.40	0.00 56.60 51.70	10.00 13.50 20.10	1.89 9.27 4.02	3.44 3.02	3.45 6.84 5.31	1.35 6.96 3.66	0.00 > 120.00 > 290.00 >	1.38	39.90	30.40 >	159.00
	Diff.	1.444 1.130 1.147	1.282 1.376 1.147	1.081 1.058 1.417	1.081 1.058 0.936	1.071 1.047 0.936	1.071	1.037 0.944 1.018	0.991	0.842	0.997	0.777	0.971
	==	YEA 269 2PP	YEC 26R 2PP	YEI 26T 2PR	YE1 26T 2PS	YE. 266. 275. 275. 275. 275. 275. 275. 275. 275	7E.J	YEK 26V 2PT	7.11 264 2PU		13K 13K 13K		Y7F YU0
	Test	08/31/90 10/03/90 10/25/90	08/31/90 10/03/90 10/25/90	08/31/90 10/03/90 10/25/90	08/31/90 10/03/90 10/25/90	08/31/90 10/03/90 10/25/90	08/31/90 10/03/90 10/25/90	08/31/90 10/03/90 10/25/90	09/05/90 10/03/90 10/25/90	08/16/90 09/13/90	08/16/90 09/13/90 08/16/90	08/16/90	08/22/90
	Ship-	222	222	222	222	222	222	222	222	22	22 22	22	22
	AVS No.	880 880 880 880	222	2082 2083 2083	7987	7085 7085 7085	7086 7086 7086	7087 7087 7087	7092 7092 7092	7319	7321 7321 7332 7332	7354	57.57 57.57

Table 9 (Cont'd)

	∀ ∪⊢	+ +	• •	+ +	+ +	+ +	. +	• •	• •	+ +	. +		* *	+ +	+ .	. +	• •	
	ASSAY	EE	FF	EE	FF	FF	FF	FE	EE	FE	ĒĒ	FF	ĒĒ	ĒĒ	ĒĒ	FF	EE	
	Į.	0.00	5.38 5.78	1.19	5.55	5.78	1.66	11.62	3.65	33.12	8.4	0.70	0.31	18.43	9.0	9.16	0.00	
	8	0.00	1.05 >	1.63	3.03 >	1.38 >	0.00	2.74	.34	55.49 >	1.25 \$	0.00	0.00	1.27	0.00	1.71	0.00	
	A1 95	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.0	0.00	0.00	0.00	0.00	0.00	0.00	0.00	
	36 21	320.00	1000.00 3200.00	961.00 3200.00	320.00 1000.00	956.00 962.00	320.00	320.00 320.00	320.00 957.00	320.00 313.00	1000.00 3200.00	320.00	320.00 883.00	320.00 320.00	0.95 3200.00	320.00 3200.00	320.00	
	10 95	0.00	0.00	0.00	0.00	9.0	0.00	0.00	9.00	9.0	0.00	0.00	0.00	0.00	302.00	0.00	, 00.00 0.00	
	A1 50	5.21	1.05	2.38	0.00	2.28	0.00	4.15	0.00	106.01	0.00	0.00	0.00	90.00	0.00	0.00	3.27	
	10 50	320.00	1000.00 >	614.00	283.00	558.00 618.00	320.00 595.00	82.20 58.20	320.00 570.00	206.00 158.00	835.00 3200.00 >	84.30 29.50	195.00 213.00	320.00 > 320.00	0.50	3200.00	320.00	
	10 50	61.40	955.00 × 0.00 ×	257.00	0.00	245.00	320.00	19.80	0.00 \$	1.94 2.39	0.00	0.00 84.10	100.00	6.34 0.00	179.00 0.00	0.00 >	0.00 >	
	A1 25	2.91	3.44	2.99	2.49	2.16	0.00	¥.5.	0.00	107.86 72.80	0.00	0.00	0.77	3.96	0.00	0.00	3.42	
	10 25	100.00	1000.0 > 3200.0	420.00	192.00	338.00 427.00	190.00 393.00	54.30	320.00	108.00 >	578.00	58.20 18.90	37.20 85.40	8.06	0.25	> 320.00 > 3200.0 >	320.00	
	10 25	34.30	290.00 \$	140.00	76.90 58.50	156.00	0.00	11.20	0.00 >	1.00	0.00	35.50	48.10	2.03	134.00	0.00 >	0.00 >	
	Diff.	0.966	0.911	0.911	0.974	0.856	0.814	0.662	0.630	1.304	1.090	1.067	0.937	1.352	1.418	1.066	0.863	
į	<u> </u>	0 Y7G	0 Y7H	0 Y7H	0 Y7X 0 21T	NZY 0	0 YAH 0 223	0 YC9	0 YCA 0 Z6M	O YUK	0 YX1	0 YX7	O YYN	O ZNG O ZYR	SYS 0	UYZ 0	0 21J	
	Test	08/22/90 Y7G 09/19/90 YUP	08/22/90 Y7H 09/19/90 YUQ	08/22/90	08/22/90 09/27/90	08/22/90	08/23/90	08/30/90 10/03/90	08/30/90 YCA 10/03/90 Z6M	09/20/90	09/20/90 YX1 10/25/90 ZPM	09/20/90 YX7 10/31/90 2UD	09/25/90	10/24/90	10/24/90	10/24/90	10/16/90 21J 12/05/90 1MK	
	Ship-	22	22	22	22	22	22	22	22	RR	RR	RR	RR	22	22	× 12	25	
	No.	25 25 25 25	757 757	37.87. 37.85	7383	7390	7424	7433	74.34	7438	7445	7461	2472	300	7906	7910 7910	7945	

Southern Research Institute

Table 9 (Cont'd)

Confirmatory Assays for Compounds Active Against Yellow Fever Virus

4 U F	+ 1
ASSAY	TIM
TAI	2.94
ïs	0.11 \$
VI 95	1.07
10 %	3000.00 > 3200.00 > 0.00 2650.00
10 %	3000.00 ,
A1 50	0.16
(C 50	271.00 671.00
10 50	1650.00
AI 25	0.00
TC 25	176.00 397.00
10 25	1190.00
Diff.	0.702
14	24V
Test	12/20/90
Ship-	22
AVS NO.	8519 8519

This value is a virus rating (VR) rather than a TAI. The VR is a measurement of selective antiviral activity that takes into account the degree of inhibition of virus-induced CPE and the degree of cytotoxicity produced by the test compound similar to TAI. TAI is more accurate with MTI measurements.

ptical densities.
optical
control
the virus control
2
Il control
n the ce
e difference in
the diff
ntial is
The differential
The
DIFRNTL =

ion (kg/ml) that inhibited viral CPE by 25%, 50% or 95% calculated by using	
(Viral) inhibitory concentration 25%, 50% and 95% = The drug concentrati	a regression analysis for semilog curve fitting.
H	
1C25.50.95	

by 25%, 50% or 95%.	th 25%, 50% or 95% reduction val
(Cell) toxicity concentration 25%, 50 % and 95 % = The drug concentration (μ g/ml) that reduced cell viability by 25 %, 50 % or 95 %.	Antiviral Index = A single point ration of the antiviral and anticellular effect of the compound, calculated with 25%, 50% or 95% reduction values
x_s 50% and 95% = The drug concentration	ration of the antiviral and anticellule
(Cell) toxicity concentration 25	Antiviral Index = A single point
H	н
TC25.50.95	AI 25.50.95

(calculated by dividing the TC _{25,50,95} by the IC _{25,50,95}).	Selectivity Index = A ratio calculated by dividing the IC ₂₅ by the IC ₅₀ (based upon 6 one-half-log ₁₀ dilutions, µg/ml, the maximum scale is 0-320).
6.00	н
4	

Total Antiviral Index = The area between the cytotoxicity and the antiviral curves (based upon a scale of 0-100%). I

Activity = A "+" denotes a test that produced ≥25% reduction in CPE. A "-" denotes an inactive test (i.e. ≤25% reduction in CPE.

ACT

4.1.4 Japanese Encephalitis Virus (JE):

The number of single drug tests carried out against JE during this contract period is summarized in yearly increments in Figure 18. During this five-year period two main in vitro antiviral assay protocols were implemented:

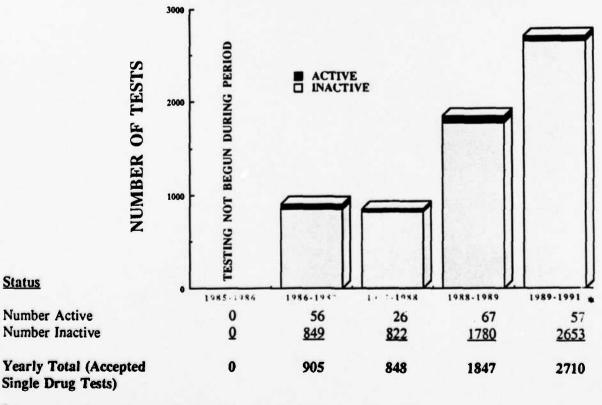
- A standard CPE inhibition assay by virus rating (VR) (Annual Report, December 15, 1. 1988, Section 3.2.4).
- 2. Since November, 1988, MTT based-antiviral assay format.

A total of 7873 tests were performed during this contract period using both assay types. Routine testing was changed to the MTT-assay format to improve the efficiency and quality of the primary screening program in addition to being more cost-effective. Selenazofurin (AVS-0253) was tested in each standard virus rating (VR) CPE-inhibition assay as a positive control compound. Results of these positive controls (VR tests) were used as a guideline to assess the quality of each assay.

After the testing was converted to the MTT-assay format, we performed a total of 372 control compound assays with Selenazofurin during the last 26 months of the contract period. During this time 781 tests were internal (+++) virus load, cell load, and other quality control tests. Four hundred ten (410) tests were considered unsatisfactory based on the criteria of the quality controls set during this reporting period. The rest, totaling 4557 were actual single drug MTT-assays. The total number of MTT-assays (6120) tested during the last two years represents a 224% increase (improvement) in the total testing output as compared to the total of 1937 tests performed during the first 3 years of this contract.

Out of the 6310 accepted single drug tests, 206 compounds demonstrated antiviral activity at greater than 50% reduction levels. This represents around 3.0% of the tested compounds having in vitro antiviral activity against JE-virus. The remainder, 6104 compounds (97%), were considered inactive with both assay protocols (Figure 18).

> IN VITRO PRIMARY SCREEN: NUMBER OF COMPOUNDS FOUND ACTIVE AGAINST JAPANESE ENCEPHALITIS VIRUS DURING THE CONTRACT PERIOD



Five-Year

Totals

206

6104

6310

Status

Number Active

Number Inactive

Represents 14-month period (November 15, 1989 - January 31, 1991) Figure 18

- 4.1.4.1 **IE-Quality Controls:** Two positive control compounds (Selenazofurin and 2-Thio-6-Azauridine) were used in the daily assay sets as antiviral activity quality control. The antiviral performance of the unknown compounds is compared to that of the positive control compounds. Compounds with equal to of better antiviral potency are considered active and are worthy of further *in vitro* profile studies and *in vivo* testing.
- 4.1.4.1.1 <u>Antiviral Activity of Selenazofurin vs JE Virus:</u> A summary of the antiviral and cytotoxicity performance of the primary control compound, AVS-0253 (Selenazofurin) is presented in Figure 19-A for 191 tests performed during November, 1989 through January, 1991.

Control Compound-Antiviral Performance: Selenazofurin (AVS-0253) has been the sole control compound against JE in these MTT-assay screens. The mean and median antiviral inhibition and cytotoxicity patterns of the positive control drug (Selenazofurin) are illustrated in Figure 19-A.

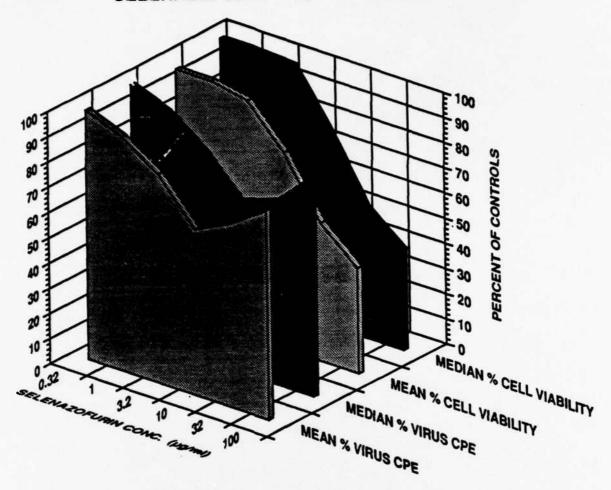
The 191 control tests performed with Selenazofurin gave a mean Total Antiviral Index (TAI) of 7.60% (SD \pm 7.60) and the median value was 5.07%. The TAI measures the overall antiviral effectiveness of the compound and it ranged from ~ 0 - 33.94% during this period. The mean Selectivity Index (SI) was only 0.39 (SD \pm 1.14) and the median SI value was 0, indicating poor antiviral selectivity for Selenazofurin and it ranged from ~ 0 - 7.17 during this period. However, the closeness of the mean and median values indicate that the present execution of the SOP is consistent and repeatable.

The mean Antiviral Index 25% (AI₂₅) value was 5.21 (SD \pm 11.5). The median AI₂₅ value was 2.11 (range 0 - 124.76). The mean Antiviral Index 50% (AI₅₀) was 1.09 (SD \pm 3.34) with a median of 0 (range 0 - 25.40). This indicates that Selenazofurin does not consistently reach 50% antiviral reduction levels. The mean Antiviral Index 95% (AI₉₅) was not attainable with Selenazofurin versus Japanese Encephalitis Virus.

The mean Antiviral Inhibitory Concentration 25% (IC₂₅) was 7.15 μ g/ml (SD \pm 8.33). The median IC₂₅ value was 5.96 μ g/ml (range = 0 - 100 μ g/ml). The mean Antiviral Inhibitory Concentration 50% (IC₅₀) was 2.84 μ g/ml (SD \pm 8.33). The median IC₅₀ value was 0 μ g/ml (range = 0 - 43.1 μ g/ml). This discrepancy indicates that the control compound Selenazofurin does not consistently reach 50% reduction levels. The mean Antiviral Inhibitory Concentration 95% (IC₉₅) could not be attained with Selenazofurin versus Japanese Encephalitis Virus.

The average maximum antiviral inhibitory level of 191 Selenazofurin tests (Figure 19-A) was reached at 10 μ g/ml of the compound with 35% antiviral effect. Further increase of the drug concentration does not improve its antiviral activity. Maximum antiviral effect (~38%) was found with a simultaneous ~25% cytotoxic suppression. Above 10 μ g/ml concentration the antiviral protection levels off to ~15% reduction level at 100 μ g/ml, while simultaneously the Selenazofurin becomes maximally toxic (~60%).

SELENAZOFURIN - VS - JE VIRUS



CONCENTRATION (µg/ml)

% Viral CPE	% Cell Viability
/V	70 Con Viability

Conc.(µg/ml)	0.32	1	3.2	10	32	100	0.32	1	3.2	10	32	100
Meau	99	93	83	66	73	83	97	97	94	78	55	42
Median	100	94	83	68	73	87	100	100	99	77	52	40
Std. Dev.	0.02	0.06	0.08	0.10	0.15	0.15	0.05	0.04	0.08	0.14	0.14	0.09

Figure 19-A

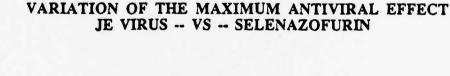
Average Antiviral and Cytotoxicity Values for 191 Positive Control Compound Tests

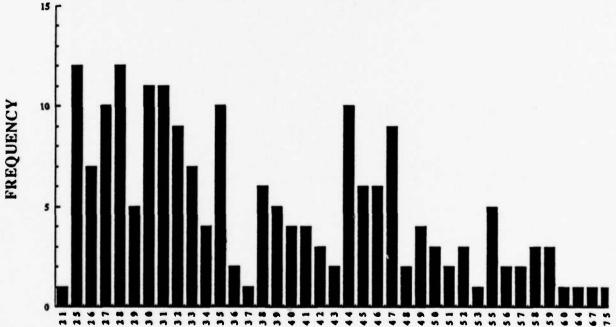
4.1.4.1.2 Maximum Antiviral Effect of Selenazofurin vs JE Virus: Since the metabolic activity of the cells was an unknown function during the testing period, it was monitored indirectly by measuring the maximum antiviral effect of the control compound Selenazofurin. This demonstrated the amount of infectious virus that was produced by the cells (Maximum Percent CPE).

A bar graph scatter plot (Figure 20-A) depicts the distribution of the maximum antiviral reduction values of all 191 control compound assays for Selenazofurin. The results indicate that the average maximum antiviral reduction obtained with the present SOP is around 38% (SD ± 10.50) reduction levels. The maximum reduction levels vary from 21 - 75% but remain quite consistently around the median of 35%. The assay control values give a relatively broad, bell-shaped distribution curve. This indicates quite a consistent day-to-day performance of the control compound in the JE-MTT assay.

During this period the positive control compound performance criteria for Selenazofurin versus the JE virus was set at 25% reduction level. Assays in which Selenazofurin did not meet this accepted quality control level (>25%) were rejected (i.e., 410 unsatisfactory tests).

Since Selenazofurin is only marginally active against JE virus, better quality control compounds are needed. Howev, regardless of the poor performance of the JE quality control drug Selenazofurin, around 142 different compounds have equal or better antiviral activity against JE virus than AVS-0253. Some of these could certainly be used as a better in vitro antiviral control compound in this large-scale antiviral screening program.





PERCENT CPE REDUCTION

Figure 20-A Maximum Antiviral CPE Reduction (%). Summary of 191 Control Tests.

4.1.4.1.3 Cellular Cytotoxicity of Selenazofurin vs JE Virus:

<u>JE-Control Compound-Cytotoxicity Performance:</u> The 191 cytotoxicity values of the positive control compound Selenazofurin are also very consistent. The mean cell Toxic Concentration 25% (TC₂₅) was 18.76 μ g/ml (SD \pm 18.19) and the median was 11.80 μ g/ml (range of 2.29 - > 100 μ g/ml). The mean cell Toxic Concentration 50% (TC₅₀) value was 57.54 μ g/ml (SD \pm 38.84) and the median was 47.10 μ g/ml (range of 7.59 - 311 μ g/ml). The mean cell Toxic Concentration 95% (TC₉₅) value cannot be attained with Selenazofurin versus Japanese Encephalitis Virus.

As can be seen from Figure 19-A, the toxicity starts to become measurable above the concentration of 3.2 μ g/ml and the maximum toxicity has not been reached at 100 μ g/ml.

When the cytotoxicity reaches around 20% (10 μ g/ml), the control compound (Selenazofurin) loses its maximum antiviral effect (35%). Above 10 μ g/ml the antiviral protection of Selenazofurin starts to decrease down to (~15%), Selenazofurin becomes maximally toxic at 100 μ g/ml concentration. The highest Selenazofurin concentration tested in these assays was 100 μ g/ml.

Selenazofurin has a definite cytotoxic suppression on cellular metabolism and growth. The TC_{25} and TC_{50} toxicity can be achieved with relative consistency at 100 μ g/ml.

4.1.4.1.4. JE-Assay Plate Quality Controls: Cell Load and Virus Load Parameters

(Selenazofurin): The MTT assay is fundamentally dependent upon the quality of the assay plates. Our large-scale antiviral testing is dependent upon the uniformity of the test plates produced for the daily assays. Equal numbers of cell load and virus load as well as the consistent performance of the reagents used daily was monitored. A sample of the plate variation control for the period of November, 1989 through January, 1991, is presented in Figures 21-A, 22-A and 23-A.

JE-Control Compound-Cell Load Performance: A bar graph scatter plot of the mean cell control (O.D. reading) of 191 control assays is plotted in Figure 21-A. The results indicate that the cell O.D. readings reached a mean 1.160 (SD \pm 0.160) with a median of 1.150 (range of 0.770 - 1.630). This indicates that a uniform and equal number (18,000 cells/well) of cells are being loaded into every well in the 96-well plate during the day-to-day operation. The cells reduced MTT to formazan giving maximum blue color uniformly and consistently.

<u>IE-Control Compound-Virus Load Performance</u>: A bar graph scatter plot of the mean virus load O.D. readings of the 191 control assays is presented in Figure 22-A. The results indicate that the average load O.D. reading is $0.310 \, (SD \pm 0.130)$ with a median of 0.330 (range of 0.040 - 0.640). This demonstrates that a good cell destruction is taking place and a uniform load of virus (32 TCID₅₀) is administered on the cell monolayer with very consistent viral CPE results.

JE-Control Compound-Assay Differential Performance: A bar graph scatter plot of the mean O.D. differential values of the 191 control assays is provided in Figure 23-A. The results indicate that the average differential O.D. reading is 0.850 (SD \pm 0.135) with a median of 0.848 (range 0.552 - 1.200). The single bell-shaped curve is reasonably sharp and uniform. This reflects that the assays are executed consistently and are repeatable during day-to-day operation with close to 85% measurement accuracy.

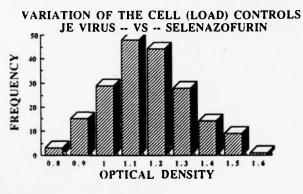


Figure 21-A

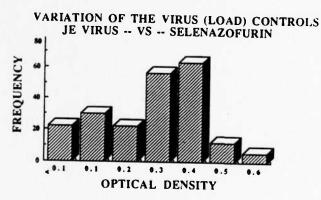


Figure 22-A

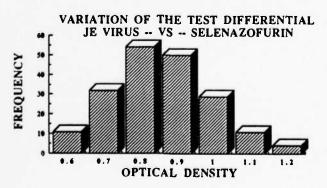


Figure 23-A

4.1.4.1 JE-Quality Controls:

4.1.4.1.1 Antiviral Activity of 2-Thio-6-Azauridine vs JE Virus: A summary of the antiviral and cytotoxicity performance of the second control compound, AVS-6724 (2-Thio-6-Azauridine) is presented in Figure 19-B for 46 tests performed during November, 1989 through January, 1991.

Second Control Compound-Antiviral Performance: 2-Thio-6-Azauridine (AVS-6724) was tested as a possible second control compound against JE in these MTT-assay screens. The mean and median antiviral inhibition and cytotoxicity patterns of this second positive control drug are illustrated in Figure 19-B.

The 46 control tests performed with 2-Thio-6-Azauridine gave a mean Total Antiviral Index (TAI) of 24.5% (SD \pm 12.20) and the median value was 20.20%. The TAI measures the overall antiviral effectiveness of the compound and it ranged from \sim 5.64 - 48.49% during this period. The mean Selectivity Index (SI) was only 8.00 (SD \pm 7.60) and the median SI value was 4.60, indicating moderate antiviral selectivity for 2-Thio-6-Azauridine and it ranged from \sim 0 - 27.27 during this period. However, the closeness of the mean and median values indicate that the present execution of the SOP is consistent and repeatable.

The mean Antiviral Index 25% (AI₂₅) value was 16.50 (SD \pm 12.30). The median AI₂₅ value was 11.70 (range 2.84 - 46.73). The mean Antiviral Index 50% (AI₅₀) was 18.50 (SD \pm 15.20) with a median of 13.60 (range 0 - 56.47). This indicates that 2-Thio-6-Azauridine does not consistently reach 50% antiviral reduction levels. The mean Antiviral Index 95% (AI₉₅) was not attainable with 2-Thio-6-Azauridine versus Japanese Encephalitis Virus.

The mean Antiviral Inhibitory Concentration 25% (IC₂₅) was 3.44 μ g/ml (SD \pm 3.20). The median IC₂₅ value was 2.50 μ g/ml (range = 1 - 21.40 μ g/ml). The mean Antiviral Inhibitory Concentration 50% (IC₅₀) was 6.67 μ g/ml (SD \pm 6.20). The median IC₅₀ value was 5.03 μ g/ml (range = 0 - 26.10 μ g/ml). This discrepancy indicates that the control compound 2-Thio-6-Azauridine does not consistently reach 50% reduction levels. The mean Antiviral Inhibitory Concentration 95% (IC₉₅) could not be attained with 2-Thio-6-Azauridine versus Japanese Encephalitis Virus.

The average maximum antiviral inhibitory level of 46 2-Thio-6-Azauridine tests (Figure 19-B) was reached at 10 μ g/ml of the compound with 60% antiviral effect. Further increase of the drug concentration does not improve its antiviral activity. Maximum antiviral effect (~70%) was found with a simultaneous ~10% cytotoxic suppression. Above 32 μ g/ml concentration the antiviral protection levels off to ~10% reduction level at 320 μ g/ml, while simultaneously the 2-Thio-6-Azauridine becomes maximally toxic (~75%).

2-THIO-6-AZAURIDINE - VS - JE VIRUS

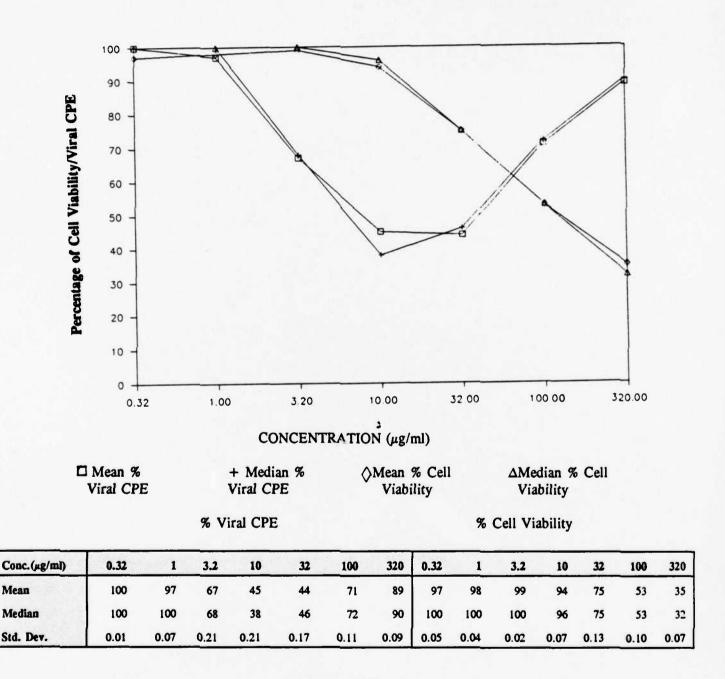


Figure 19-B
Average Antiviral and Cytotoxicity Values for 46 Positive Control Compound Tests

4.1.4.1.2 <u>Maximum Antiviral Effect of 2-Thio-6-Azauridine vs JE Virus:</u> Since the metabolic activity of the cells was an unknown function during the testing period, it was monitored indirectly by measuring the maximum antiviral effect of the control compound 2-Thio-6-Azauridine. This demonstrated the amount of infectious virus produced by the cells (Maximum Percent CPE).

A bar graph scatter plot (Figure 20-B) depicts the distribution of the maximum antiviral reduction values of all 46 control compound assays for 2-Thio-6-Azauridine. The results indicate that the average maximum antiviral reduction obtained with the present SOP is around 70% (SD \pm 17.50) reduction levels. The maximum reduction levels vary from 46 - 100% but remain quite consistently around the median of 64%. The assay control values give a relatively broad, bell-shaped distribution curve. This indicates quite a consistent day-to-day performance of the control compound in the JE-MTT assay.

Recommendations:

Based upon the data obtained in parallel studies with Selenazofurin, we recommend that 2-Thio-6-Azauridine (AVS #6724) be used as a second control compound against JE virus. It's overall performance is much better than the present control, Selenazofurin. It is readily available from Sigma Chemical Company, it is inexpensive and works as effectively at low drug concentrations as Selenazofurin.

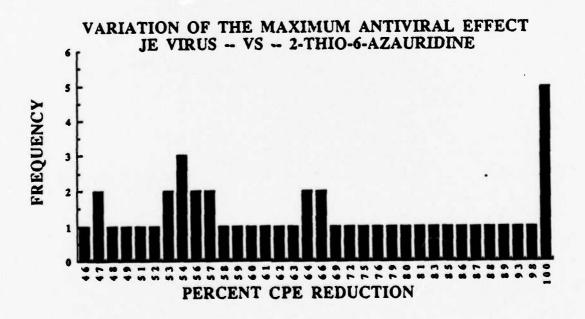


Figure 20-B

Maximum Antiviral CPE Reduction (%).

Summary of 46 Control Tests.

4.1.4.1.3 Cellular Cytotoxicity of 2-Thio-6-Azauridine vs JE Virus:

<u>JE-Control Compound-Cytotoxicity Performance</u>: The 46 cytotoxicity values of the positive control compound 2-Thio-6-Azauridine are also very consistent. The mean cell **Toxic Concentration** 25% (TC₂₅) was 40.20 μg/ml (SD \pm 22.50) and the median was 33.40 μg/ml (range of 9.50 - 100 μg/ml). The mean cell **Toxic Concentration** 50% (TC₅₀) value was 98.20 μg/ml (SD \pm 41.10) and the median was 100 μg/ml (range of 41.70 - 320 μg/ml). The mean cell **Toxic Concentration** 95% (TC₉₅) value cannot consistently be attained with 2-Thio-6-Azauridine versus Japanese Encephalitis Virus.

As can be seen from Figure 19-B, the toxicity starts to become measurable above the concentration of 10 μ g/ml and the maximum toxicity has not been reached at 320 μ g/ml.

When the cytotoxicity reaches around 25% (32 μ g/ml), the control compound (2-Thio-6-Azauridine) loses its maximum antiviral effect (70%). Above 32 μ g/ml the antiviral protection of 2-Thio-6-Azauridine starts to decrease down to (~10%), becomes maximally toxic at 320 μ g/ml concentration. The highest 2-Thio-6-Azauridine concentration tested in these assays was 320 μ g/ml.

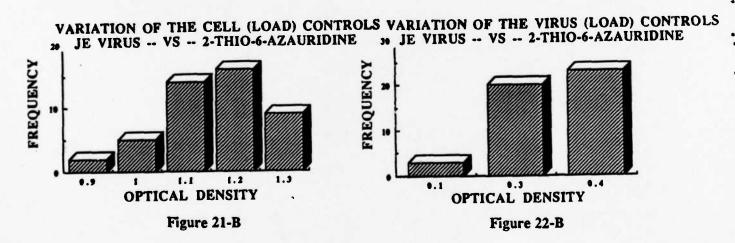
2-Thio-6-Azauridine has a definite cytotoxic suppression on cellular metabolism and growth. The TC_{25} and TC_{50} toxicity can be achieved with relative consistency at 100 μ g/ml.

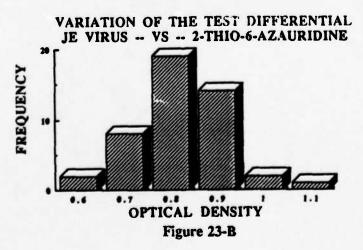
4.1.4.1.4. IE-Assay Plate Quality Controls: Cell Load and Virus Lead Parameters (2-Thio-6-Azauridine): The MTT assay is fundamentally dependent upon the quality of the assay plates. Our large-scale antiviral testing is dependent upon the uniformity of the test plates produced for the daily assays. Equal numbers of cell load and virus load as well as the consistent performance of the reagents used daily was monitored. A sample of the plate variation control for the period of November, 1989 through January, 1991 is presented in Figures 21-B, 22-B, and 23-B.

<u>IE-Control Compound-Cell Load Performance</u>: A bar graph scatter plot of the mean cell control (O.D. reading) of 46 control assays is plotted in Figure 21-B. The results indicate that the cell O.D. readings reached a mean 1.160 (SD \pm 0.100) with a median of 1.160 (range of 0.930 - 1.350). This indicates that a uniform and equal number (18,000 cells/well) of cells are being loaded into every well in the 96-well plate during the day-to-day operation. The cells reduced MTT to formazan giving maximum blue color uniformly and consistently.

IE-Control Compound-Virus Load Performance: A bar graph scatter plot of the mean virus load O.D. readings of the 46 control assays is presented in Figure 22-B. The results indicate that the average load O.D. reading is 0.330 (SD \pm 0.070) with a median of 0.350 (range of 0.080 - 0.440). This demonstrates that a good cell destruction is taking place and a uniform load of virus (32 TCID₅₀) is administered on the cell monolayer with very consistent viral CPE results.

<u>IE-Control Compound-Assay Differential Performance</u>: A bar graph scatter plot of the mean O.D. differential values of the 46 control assays is provided in Figure 23-B. The results indicate that the average differential O.D. reading is 0.820 (SD \pm 0.100) with a median of 0.820 (range 0.575 - 1.117). The single bell-shaped curve is reasonably sharp and uniform. This reflects that the assays are executed consistently and are repeatable during day-to-day operation with close to 82% measurement accuracy.





4.1.4.2 JE-Antiviral Activity Results:

New Drugs with 95% Antiviral Reduction Levels: Out of the 6310 actual single drug tests, 45 new compounds demonstrated excellent antiviral activity, having antiviral reduction values of equal to or better than 95%. This represents around 0.7% of the test compounds being active at this excellent reduction level. These compounds are summarized in Table 10 according to the highest Total Antiviral Index (TAI). Compounds AVS-5580 demonstrated the greatest in vitro promise, having a TAI of 96% and Selectivity Index (SI) of >313. The next 23 compounds demonstrated good antiviral activity with TAI's that ranged from 25 - 62 and SI values that ranged from <1 - 41. Twenty-one other compounds demonstrated moderate antiviral activity, having TAI's that ranged from 9 - 24% and SI's from 2 - 5.

It is worthwhile to note that compounds received in shipment number 62 were mostly colored (Table 10). Therefore those compounds appearing in the 95% active category from shipment number 62 should be interpreted with caution, since colored compounds create false positive readings with the MTT assay.

Table 10

AVS Compounds Active Against Japanese Encephalitis Virus (JE) at AI95 Level

	AVS	Ship-	Test	Diff-									
Virus		ment#		rntl.		IC 95		TC 95		AI 95	SI		TAI
		mette.	Date	THEE.		10 ,5		10 93	•	WI 33	31		INI
JE	5580	54	05/03/89	0 486	-	320.00	>	100000	•	313.00 >	313 00	•	95 79
	4855	48	09/13/89		•	28.80		320.00		11.12 >			
	4796	46	09/13/89			3.05		100.00		32.78	40.51		
	2631	65	07/06/90			71.50		320.00		4.47 >	36.90		
	3802	67	07/12/90			0.93		10.00		10.75 >			
	2318	13	08/30/89			0.87		10.00		11.45	24.21		
	0148	67	06/21/90			0.89		3.20		3.59 >	10.00		
	2563	48	02/06/89			0.89		3.20		3.59 >			
	2980	48	03/10/89			0.73		3.20		4.39	12.00		40.06
	4527	47	01/31/89			7.72		94.60		12.30	11.20		
	2811	48	02/06/89			0.09		0.91		10.40	11.00		
	0206	67	06/21/90					1000.00		10.74	12.94		36.80
	8353	75	11/29/90			97.80		320.00		3.27 >			
	4592	48	02/28/89			26.50		100.00		3.77	7.19		32.32
	5138	57	07/11/89			94.30		320.00		3.77	0.16		
	4609	48	02/28/89			0.03		0.10		3.22 >			
	0361	48	02/26/89			0.03		0.30			4.30		
	6214	62	11/01/90					1000.00	_	10.40 3.40	3.82		
7.77	6195	62				292.00	-	966.00	-	3.40			29.00
	6219	62	10/31/90 11/01/90			282.00	_	320.00	_	1.13 >			
	0360							1.00				>	
7.77		2	08/23/89			0.09	>		>	10.84	5.64		27.74
100	4113	39 65	05/24/89			0.28 9.16		2.85		10.10	4.01		
75.51	4277		07/06/90					93.80		10.24	4.12		
7 7		42	09/13/89			2.92		10.00		3.42	4.39		
	4278 5539	42 56	09/13/89			9.40	>	32.00	>	3.40	5.15		
		-	08/09/89			3.20		94.30		29.48	3.94	>	
	5121	56	08/08/89			91.50		309.00		3.38	3.78		22.70
	4590	48	02/28/89			0.94		28.70	_	30.60	4.11		
	4074	48	03/10/89			0.03		0.03		1.09 >			
	5484	53	12/12/89				>	1000.00	>	1.09 >			
	6225	62	11/01/90			301.00		964.00		3.21	2.70		
	6207	62	10/31/90			302.00		966.00	_	3.20	2.74	>	
	2812	48	02/06/89			0.01	>	0.01	>	1.06 >			18.14
	8352	75	11/29/90			3.06		32.00	_	10.45	3.58		
7.0	6946	69	07/26/90			301.00	>	320.00	>	1.06 >		>	
	5405	66	07/06/90			0.30		9.07		29.84	3.09		16.46
	6203	62	01/11/90			94.50		320.00		3.39	3.08		16.40
7.00	1337	33	08/29/89			29.40	>	32.00	>	1.09 >	2.30		
	6986	68	08/09/90			302.00		973.00		3.22	2.75	>	
	6204	62	01/11/90			30.20		96.60		3.20	2.74		14.73
. 3	4070	48	03/10/89			2.99		3.20		1.07 >	1.98		
	6199	62	01/11/90			94.50		320.00		3.39	2.80		_
	4280	42	09/13/89			9.29	>	10.00	>	1.08 >	2.10		
	6196	62	10/31/90			302.00		952.00		3.15			10.13
JE	7445	73	10/25/90	0.769		3020.00	>	3200.00	>	1.06 >	1.79	>	8.72

New Drugs with 50% Antiviral Reduction Levels: Out of the 6310 actual single drug tests, 84 new compounds demonstrated good antiviral activity, having antiviral reduction values equal to or better than 50%. This represents around 3% of the test compounds being active at this good antiviral reduction levels. These compounds are summarized in Table 11 according to the highest Total Antiviral Index (TAI). AVS-0217, 5998 and 1645 demonstrated the best TAI is of $\sim 40\%$ and SI is that ranged from 11 - 18. The next six compounds demonstrated moderate antiviral activity, having TAI's that ranged from 25 - 29% and SI's from 4 - 9. The rest (74) compounds showed marginal antiviral activity with TAI's that ranged from < 1 - 23 and SI's that ranged from < 1 - 4.

It is worthwhile to note (Table 11) that compounds received in shipment number 62 were mostly colored. Therefore those compounds appearing in the 50% active category from shipment number 62 should be interpreted with caution, since colored compounds create false positive readings with the MTT assay.

Table 11 AVS Compounds Active Against Japanese Encephalitis Virus (JE) at ${\rm AI}_{50}$ Level

*** 1. 2	AVS	Ship-		Diff-							
Virus	NO.	ment#	Date	rnt1.	IC 50		TC 50		AI 50	SI TAI	
JE	0217	33	09/27/89	0.786	28.10	>	320.00	>	11.39 >	11.39 > 41.60	
	5998	61	02/01/90		1.69		32.00		18.94	18.94 > 41.36	
	1645	33	08/29/89		1.79	>	32.00	>	17.89	13.28 > 40.13	
	2979	48	03/10/89		2.11		21.20		10.10	5.91 28.50	
	2503 0111	21 42	09/06/89 05/10/89		0.43 48.80		9.43 320.00		22.06 6.56	6.78 > 28.10 4.45 27.93	
	4785	46	09/13/89	0.490	1.56		22.10		14.16	5.94 > 27.84	
	5997	61	11/07/89		3.51		75.30		21.47	5.23 > 25.75	
JE	0646	2	08/23/89		0.21		3.15		15.21	8.72 > 25.70	
	1019	28	03/06/90		2.57		78.8 0		30.61	5.79 24.72	
	0094	1	09/27/89		31.00	>	100.00	>	3.23 >	3.23 > 23.29	
	5532	56	08/09/89		172.00		663.00		3.85	2.85 > 21.79	
	4871 4978	61 27	11/29/89 12/05/89		153.00	>	1000.00 857.00	>	5.09 5.59	3.85 21.64 3.64 21.45	
	5515	53	04/18/89		11.20		58.40		5.20	3.04 21.27	
	3964	33	03/21/89	0.815	6.61		32.30		4.89	2.43 19.74	
JE	5695	57	08/22/89		21.70	>	32.00	>	1.47 >	1.47 > 18.09	
	6413	66	07/06/90	0.718	32.00		158.00		4.94	2.66 > 17.63	
	5503	53	04/18/89		25.60		154.00		6.03	2.98 17.26	
	5601		02/08/90		1430.00	>		>	2.23	2.23 > 17.12 7.45 17.05	
	2275 5497	12 53	08/30/89 12/20/89	0.803	22.40 221.00		234.00 767.00		10.43 3.48	2.46 16.05	
	6942	69	10/24/90		95.90		205.00		2.14	1.55 > 16.04	
	5142	57	08/16/89		56.10		210.00		3.74	2.76 15.95	
	1644	64	05/03/90		139.00		813.00		5.85	0.47 15.25	
	7911	75	11/07/90		238.00		938.00		3.95	2.65 > 15.09	
	4452	44	07/07/89		68.40	_	277.00		4.05	2.61 14.99	
	0084 7949	1 75	08/23/89 12/12/90		49.20 235.00	>	100.00 950.00	>	2.03 > 4.04	2.03 > 14.89 2.70 14.83	
	1381	45	02/13/90		24.10		81.30		3.38	2.35 > 14.76	
	7321	70	08/15/90	0.753	20.70		86.80		4.19	2.87 > 14.47	
	0124	64	03/08/90		226.00	>	320.00	>	1.42 >	1.42 > 14.11	
	4739	44	07/07/89		19.00		66.00		3.47	2.54 > 13.96	
	2506	64	03/13/90		10.00		68.00		6.80	1.99 13.02	
10.70	6206 0347	62 41	01/11/90 05/10/89	0.776	60.30 2.08		210.00 8.55		3.48 4.10	2.57 12.46 2.37 12.33	
	6228	62	01/16/90	1.007	62.60		210.00		3.36	2.48 12.33	
	6227	62	01/16/90		80.00		210.00		2.63	1.94 > 12.09	
	4992	61	11/29/89	1.017	249.00		660.00		2.65	1.96 11.74	
	5906	61	10/31/89		2.12		15.00		7.08	2.00 11.52	
	5495	53	04/18/89		1.42		5.75		4.04	1.85 > 11.51	
	8263	76	12/12/90	0.741	66.30		232.00		3.50	2.31 11.47	
	5035 4098	48 37	10/10/89 02/24/89		1.72		6.26 0.00		3.65 1.37 >	0.19 11.29 1.37 > 11.24	
	5483	66	07/18/90		0.07		0.26		3.64	2.52 > 10.78	
	0083	64	03/08/90		3.11		10.40		3.36	2.15 > 10.68	
JE	2034	56	08/08/89	0.829	0.65		2.43		3.72	2.62 10.63	
	5538	56	08/09/89	0.866	207.00		587.00		2.84	1.80 > 10.49	
	6234		02/08/90		2670.00	>		>	1.20 >	1.20 > 10.07	
	6477 0230	66 1	07/11/90 08/23/89		22.70 94.00		72.50		3.19 1.06 >	2.30 > 9.27 1.06 > 9.02	
	4769	44	11/08/88		77.80		169.00		2.18	1.22 > 8.82	
	8499	75	11/29/90		221.00	>	320.00	>	1.45 >	1.45 > 8.53	
JE	4768	44	11/08/88		75.70		165.00		2.18	1.18 8.52	
	5058	48	02/27/89		310.00	>	320.00	>	1.03 >	1.03 > 8.39	
	7087	72	08/31/90		8.11		30.00		3.70	1.08 > 8.05	
	5905	61 62	10/31/89		7.06		19.10		2.70	1.78 > 8.02	
	6202 7083	72	01/11/90 10/03/90		219.00 83.30		320.00 191.00		1.46 > 2.29	1.46 > 7.56 1.51 > 7.55	
	7910	75	11/07/90		2220.00	>		>	1.44 >	1.44 > 7.46	
JE	7085	72	10/25/90	0.811	24.60		65.30		2.65	1.30 7.29	
	2992	27	09/07/89		83.30		100.00		1.20 >	1.20 > 6.27	
	5489	66	05/08/90		282.00	>	320.00	>	1.14 >	1.14 > 6.14	
JE	4984	51	10/10/89	1.087	7.52		21.20		2.81	1.78 5.98	

Table 11 (Cont'd)

	AVS	Ship-	Test	Diff-							
Virus	No.	ment#	Date	rntl.	IC 50		TC 50		AI 50	SI	TAI
JE	5535	56	06/14/89	0.967	204.00	>	320.00	>	1.57	1.44 >	5.90
JE	5053	48	10/10/89	1.029	56.60	>	100.00	>	1.77	1.37 >	5.79
JE	7319	70	08/15/90	0.776	10.00		18.40		1.84	0.84 >	5.48
JE	5485	53	12/14/89	0.864	25.80	>	32.00	>	1.24	1.10 >	5.44
	8364	76	12/13/90		222.00		580.00		2.62	0.33	5.41
JE	6369	63	03/20/90	1.018	28.30		61.40		2.17	1.49	5.16
JE	7433	70	08/30/90	1.049	26.00		73.10		2.81	1.58	5.08
JE	8374	76	12/13/90	0.903	824.00		1840.00		2.24	1.41	5.00
JE	4996	51	10/10/89	1.085	232.00	>	320.00	>	1.38	1.38 >	4.44
JE	1838	64	11/01/90		259.00		558.00		2.15	1.30	4.29
JE	5072	48	02/28/89	0.724	207.00	>	320.00	>	1.54	0.42 >	3.42
JE	8221	75	11/21/90	0.859	29.20		60.60		2.08	0.97	2.95
JE	6617	64	04/19/90		814.00	>	1000.00	>	1.23	1.12 >	2.49
JE	8372	76	12/13/90	0.889	2550.00	>	3200.00	>	1.25	1.08 >	2.29
JE	7051	72	10/03/90	1.019	749.00	>	1000.00	>	1.34	0.63 >	2.09
JE	7461	73	10/31/90	0.866	90.20		158.00		1.76	0.25 >	1.85
JE	2600	65	04/12/90	0.881	802.00	>	1000.00	>	1.25	0.37 >	1.75
JE	5643	57	07/06/89		230.00		282.00		1.23	0.04 >	0.68
	7375	70	08/23/90		313.00			>	1.02	0.30 >	0.55
	7045	69	08/02/90		288.00	>	320.00	>	1.11	0.60 >	0.52

New Drugs with 25% Antiviral Reduction Levels: Of the 6310 actual single drug tests, 180 new compounds demonstrated moderate antiviral activity, having antiviral reduction values equal to or better than 25%. This represents around 2.9% of the test compounds being active at this marginal antiviral reduction level.

In general, when compared to the 95% and 50% antiviral activity categories, the compounds in this (25%) category do not appear to have any significant antiviral promise and probably do not need to be presently confirmed any further.

4.1.4.3 Confirmatory Assays:

Some of the compounds were sent to us in more than one separate drug shipment. These compounds were tested more than once. Data from the confirmatory assays are summarized in Table 12. If a compound showed $\geq 50\%$ reduction in CPE during this contract period, then it was considered a candidate for confirmatory testing. The confirmatory tests are from active compounds picked up by both the VR and MTT assay testing. Out of 119 confirmatory tests, 91 compounds were confirmed active during this reporting period and the remaining 28 compounds gave conflicting results. The criteria for activity is that the confirmatory test has to show $\geq 25\%$ reduction in CPE. Failure to confirm the activity in these compounds was probably due to differences during the assay conditions:

- 1) In confirmatory assays the concentration range is adjusted to a more accurate semilog scale to maximize the TAI window.
- 2) Differences in the "differential" of the two runs can cause the compound to read positive or negative, falsely. The variability in the differential can cause false positive or negative bias to be introduced into the calculations, thus reflecting the variability in the maximum activity of the compound.
- 3) The metabolic rate, cell and virus load/well, age, and passage number of the cells may cause the above observed variability in the confirmatory results.
- 4) Problems associated with stability and storage of the compound (i.e., different lot numbers, solubility, light sensitivity, hygroscopic, etc.).
- 5) Problems associated with technical execution of large numbers of plates by different technicians.

During this reporting period the overall confirmatory rate against JE was 76%. The conflicting results should be retested at a later date based on the availability of the compound.

4.1.4.4 Recommendations of JE-Actives Based Upon the In Vitro Results with MTT Assay (Vero Cells).

Based upon the in vitro results with the MTT assay (Vero cells) we recommend the following:

- 1) Compounds with the highest TAI in the 95% activity category that have confirmed results with the exception of "colored" compounds should receive the highest priority for further profile studies and in vivo animal testing.
- 2) Compounds with the highest TAI in the 50% activity category that have confirmed results with the exception of "colored" compounds should receive the next highest priority for further profile studies and in vivo animal testing.

Table 12

Confirmatory Assays for Compounds Active Against Japanese Encephylitis Virus (JE)

< ∪ ⊢	* *	* * *	• • •	* * * * * *	* * * •	+ + +	+ + · · + +	*** ***	+ + +
Assay Type	3 3			CPE CPE CPE MIT				MTT MTT CPE CPE	
141	0.20	10.68 0.83 9.45	2.65 14.89 12.20	1.80° 2.80° 1.50° 0.00	27.93 4.54 12.14 4.38	14.11	2.10 1.80 0.00 0.00 1.10		36.80
IS	0.13	2.15 × 0.00 0.00	2.51	5.20 5.50 0.10 0.00	7.75 0.00 0.00 0.00	0.00	10.40 3.40 0.00 0.70 1.20	A A A	0.00 0.00 2.94
8	0.00	0.00	0.00	0.0.00 0.00 0.00 0.00 0.00	90000	0.00	1.00	A A	0.00 0.00 10.74
TC 95 AI	100.00	29.80 24.00 28.50	320.00 100.00 314.00	100.00 320.00 > 320.00 100.00 1000.00	320.00 1000.00 1000.00	320.00 1000.00 1000.00	320.00 > 320.00 > 100.00 320.00 320.00 320.00 320.00 320.00 320.00 320.00	^ ^	1000.00 1000.00 1000.00 v
10 % 11	0.00	0.00	0.00	320.00 × 3 0.00 × 3 0.00 × 3 0.00 × 10 0.00 × 10 0.00 × 3	0.00 0.00 0.00 0.00 0.00 0.00 0.00 0.0	0.00 \ 10	320.00 × 3 0.00 × 1 0.00 × 2 0.00 × 2 320.00 × 3	^ ^ ^ ^ ^ ^	0.00 v 10 0.00 v 10 0.00 v 10 0.00 v 10
AI 50 I	0.13	3.36	0.00 2.03 3.68	5.20 5.50 - 3 0.10 0.00 0.00	6.56 0.00 0.00	1.42	3.40 3 0.00 0.00 0.70 1.20 3	•	0.00 0.00 19.36
10 50 /	3.20	10.40 5.62 7.62	91.10 100.00 > 88.70	100.00 100.00 32.00 100.00 > 955.00	320.00 × 100.00 1000.00 879.00	320.00 > 61.60 90.80	320.00 > 320.00 > 100.00 320.00 86.00 100.00 100.00 320.	3.20 > 3.20 > 2.63	813.00 793.00 950.00
IC 50	24.00	3.11 0.00 0.00	0.00 49.20 > 24.10	19.00 × 246.00 × 31.00 × 0.00 1	48.80 × 0.00 × 0.00 × 1 × 00.0	226.00 × 0.00 0.00	31.00 94.00 > 0.00 > 121.00 83.00	0.77 > 0.32 > 0.00 - 108.00 > 178.00 - 0.00 > 0.00	0.00 0.00 49.10
AI 25	3.20	1.34	0.00 3.07 4.54	10.00 31.00 1.00 23.31 0.00	26.80 1.66 2.84 0.00	0.00	5.00 4.80 0.00 0.00 1.50	0.12 22.95 1.31 10.00 0.80	3.56 0.00 18.47
10 25 1	1.00	6.67 2.87 5.41	54.20 100.00 > 60.30	100.00 × 100.00 × 100.00 × 100.00 × 388.00 188.00	217.00 100.00 > 616.00 303.00	320.00 > 25.20 58.20	100.00 320.00 > 100.00 320.00 86.00	0.03 3.20 > 0.74 320.00 > 100.00 253.00	566.00 547.00 635.00
10 25	10.00	1.50 2.15 1.59	0.00 32.50 × 13.30	10.00 × 3.20 100.00 - 4.29 × 0.00 286.00	8.10 60.30 > 217.00 0.60	55.70 × 3.00 0.00	21.00 66.00 v 0.00 v 86.00 v	0.26 0.14 > 0.57 32.00 > 125.00 115.00	
Diff.	\$ ≨	0.686 0.914 0.911	0.959	NA NA NA 0.766 0.880 1.014	0.741 0.643 0.672 0.828	0.633	111111	0.579 0.925 NA NA ~	0.770 1.007 0.579
Test Pit Date #	01/27/87	03/08/90 UFY 05/01/90 VSO 11/01/90 2VT	05/23/89 QSP 08/23/89 RAN 06/21/90 UYW	03/09/87 01/29/88 05/04/88 09/27/89 RS9 04/10/90 V6Y	05/10/89 QOW 08/23/89 RAP 12/07/89 STG 12/07/89 STG	03/08/90 UG0 05/01/90 VS0 07/18/90 XK0	03/20/87 ···· 07/28/87 ···· 12/07/87 ···· 01/29/88 ···· 05/04/88 ····		12/05/89 SRU 12/05/89 SRN 06/21/90 WYX
Ship- Te	4	22 23 25 11, 15, 11, 12, 12, 13, 13, 13, 13, 13, 13, 13, 13, 13, 13	1 05/ 1 08/ 67 06/	1 03/ 1 01/ 1 05/ 1 09/ 65 04/	42 05/ 42 08/ 9 12/ 9 12/	\$4 03/ 69 07/	4 03/ 4 12/ 4 01/ 4 05/ 4 05/	79/	4 12/ 4 12/ 67 06/
AVS SI No. m	0079 9 0079 9	0083	7800	7600 7600 7600 7600	5555	0124 0124 0124	0136 0136 0136 0136		0206 0206 0206

(U F	+ + +	• •	+ + •	+++++		*** ****** *** ***
Assay Type	EEE	FF	THE THE	CPE CPE MITTIE	CPE CPE MITT MITT MITT MITT MITT MITT MITT MIT	
TAI	2.11	5.07 10.63	12.86 17.05 0.98	3.30° 2.50° 0.00 43.71 2.68 34.12	3.60 1.20 28.10 2.51 2.51 2.51 4.08	8.07 13.02 0.00 1.00° 1.00° 1.50° 2.32 16.57 32.03 57.59 14.21
15	0.64	0.00	0.00	14.00 6.30 0.00 24.21 v 14.30 v	5.10 6.00 6.00 7.85 7.00 0.00 0.00 0.00	24.00.00.00.00.00.00.00.00.00.00.00.00.00
A1 95	888	0.00	0.00	10.00 10.00 11.45 0.00	320.00 3.10 0.00 0.00 0.00 0.00	0.00 0.00 0.00 0.00 0.00 0.00 0.00 0.0
70 95	320.00 952.00 956.00	9.62	320.00 320.00 320.00	320.00 × 10.00 × 10.00 × 10.00 × 10.00 × 10.00	320.00 10.00 10.00 10.00 10.00 32.00	320.00 320.00 320.00 3.20 3.20 3.20 3.20
10 95	9.8.6	0.00	0.00	3.20 1.00 0.00 0.87 0.00 0.00	1.00	0.00 0.00 0.00 0.00 0.00 0.00 0.00 0.0
A1 50	1.27	3.72	0.00 10.43 0.00	14.00 20.00 0.00 36.69 19.36	5.30 1.80 0.00 22.06 0.00 12.28 0.00	5.30 6.80 0.00 1.30 1.30 1.30 0.00 0.00 36.90 31.00 0.00
TC 50	320.00 > 521.00 558.00	2.78	216.00 234.00 237.00	0.00 0.00 0.00 0.00 0.00 0.00 0.00	3.20 3.20 3.20 10.00 2.92	10.00 18.80 3.20 3.20 3.20 20.80 20.80 20.80 3.20 1.24 3.20 3.00 3
10 50	252.00 × 304.00 × 259.00	0.00	0.00	0.70 0.50 \$ 0.00 0.27 \$ 0.00	0.19 1.80 0.00 0.43 0.00 0.00	7.02 \ 0.00 \ 0.
AI 25	1.01	3.05	5.80 12.19 0.00	31.30 10.00 0.00 39.98 1.73 21.60	3.00 1.00 15.81 2.22 0.58 2.51	7.53 7.53 7.00 7.10 7.53 7.56 7.56 7.56 7.50 7.51 7.51 7.51 7.51 7.51 7.51 7.51 7.51
TC 25	160.00 247.00 338.00	1.83	106.00 167.00 162.00	10.00 ~ 3.20 ~ 1.00 6.60 0.93 7.38	3.20 × 2.90 × 1.30 × 1.30 × 1.30 × 1.30 × 1.30	10.00 1.30 1.30 1.30 1.30 1.30 1.30 1.30
10 25	159.00 162.00 161.00	0.60	18.00 13.70 0.00	0.32 0.00 0.17 0.54 0.34	0.32 1.00 0.18 0.59 0.51 0.51	6.52 6.52 6.53
Diff.	1.063 0.869 0.868	0.957	0.811 0.803 0.972	NA - NA - 0.791 0.686 0.972 0.871	NA N	0.996 1.032 0.911 NA NA 0.850 1.087 0.822 1.046 0.917 0.766 NA 1.248 <
= *	90 UIQ 90 VVN	789 08Y 789 R1U	/89 PF5 /89 RE3 /90 UGP	/87 /89 PF6 /89 RE4 /90 MGP	/87 /88 /89 PF7 /90 UGQ /90 XG7	790 VNN
Test	03/13/90 05/03/90 11/01/90	06/13/89	04/10/89 08/30/89 05/31/90	05/05/87 07/28/87 04/10/89 08/30/89 05/31/90	06/18/87 05/25/88 04/10/89 09/06/89 05/31/90 10/04/90	09/06/89 03/13/90 05/03/90 05/03/90 05/25/88 05/06/89 11/01/90 09/06/89 04/17/90 07/06/90 05/23/89
Ship- ment	333	22	53 12 67	13 13 67	22 23 23 24 26 26 26 26 26 26 26 26 26 26 26 26 26	\$25 655 458333 65 23 556 556 557 557 557 557 557 557 557 557
AVS No.	1838 1838 1838	2034	227 2275 2275	2318 2318 2318 2318 2318 2318	2503 2503 2503 2503 2503 2503 2503	2506 2506 2506 2503 2503 2503 2503 2503 2503 2503 2503

Table 12 (Cont'd)

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Assay	170	GPE	CPE	HTT	ě	ב ב	HT	HT	H	CPE	197					=	CPE	CPE	TTH	HTT	TIM	H	GE	TO TO	; <u> </u>	į	TIM	HTT	HTT	TIM	MT.	E		CPE	HTT	MI	Ħ	TTM	HIT	HT	H
	Į.	1.20	1.10	37.18		3	18.14	8.53	0.00	1.50	0 40	27 .	60.00	20.00	20.0	07.	2.30	1.10	15.31	40.09	17.45	21.71	2.30	000	2 2	3	20.53	0.28	45.81	12.64	10.74	4.70		1.50	2.68	13.87	18.93	0.00	19.27	7.52	11.27
	15	09.0	3.50	11.00 \$	8	2.40	2.57	0.00	0.00	5.10	1 20	25.0	0.0			3	16.00	1.20	2.48 >	12.00	3.37 >	3.51 >	3,10	0 0	8 6	3	4.25	0.06	20.57 >	1.30	2.43	0.89		1.50	1.32 >	1.98 >	.8	0.00	2.30 >	1.98	1.18
	A1 85	10.00	1.00	10.40	9	3	1.06	0.00	0.00	0.00	00 0		88	3 8	3 6	3	10.00	0.00	0.0	4.39	0.00	0.00	32.00	000	8 6	3	0.00	0.0	10.73 V	00.00	00	0.0	8	3.20	0.00	1.07 >	0.00	0.00	7.8	26.8 >	0.0
	8	3.20	1.00 >	0.91	620	0.036	0.01	0.26	0.32	109.00	42 00	26.40	60.10	20.00	30.00		32.00	10.00	3.20	3.20 >	10.00	3.20	32.00	6	20 00	20.03	10.00	10.00	10.00 >	160.00	32.00	9.76	5	95.00	3.20	3.20 >	45.60	0.03	0.03 >	26.80 >	10.00
	5 8	0.32	1.00 >	0.0	6	3	0.01	0.0	0.00	0.00	00.0		38	88	88	3	3.20	0.00	0.00	0.73 >	0.00	0.00	1.00	* 00 0	^ 00		0.00	0.00	0.93 >	0.00	0.00	0.0	0	10.00	0.00	2.99 >	0.0	0.00	0.03 >	9:	0.00
	A1 50	5.30	3.00	22.20	0, ,	2	2.57	0.00	0.00	16.00	A 80	72.		2.4	- - - -	3	53.00	13.00	4.62	18.20	7.26	5.15	320.00	7 30	8 8	3	52.70	0.26	20.57	2.57	68.7	2.58	3	46.00	1.31	1.98	3.66	0.00	2.30	5.%	2.6%
	TC 50	3.20	1.00	0.47	620	20.0	0.01 v	6.0	0.32	100.00	32 00 -	25.5	25.50	03.13	16.60	•	32.00 -	10.00	3.20 >	3.20 >	1.94	2.58	32.00 -		8 8	3	10.00 >	1.00	10.00 >	105.00	32.30	24.90		52.00 ×	3.20 >	3.20 >	16.40	0.03	0.03 >	2.96 ^	1.80
	10 50	0.60	0.30 >	0.02	900		0.00	0.0	0.00	6.20 -	8 40 -	27 +	. c		= =	3	- 09.0	0.80	< 69.0	0.18 >	0.27	0.50	0.10 ~	0 30 ×	8 6	3	0.19 >	3.85	< 65.0	09.07	6.61	2.0	í	c. 22 -	2.45 >	1.61 >	4.47	0.00	0.01	8.	89.0
	A1 25	1.00	10.00	32.70	9	3	4.89	4.11	0.00	10.00	2.00	/2 +	† ř	- 0	1 56	2	31.00	2.00	4.13	19.10	2.90	7.45	1.00	01.0	00	3	6.47	0.69	29.50	5.72	10.20	2.23	8	2.00	2.45	2.79	9.14	0.00	4.20	1.98 <	3.93
	52 52	0.32	1.00 -	0.24	20	20.0	0.01	0.05	0.32	32.00 -	10.00	200	2,00	20.7	6	2.5	10.00	1.00	1.72	2.10	0.0	1.76	0.32	0.01	9	3	0.81	0.54	10.00 >	53.00	16.00	8.57		3.6	3.24	3.20 >	8.51	0.03	0.03 >	1.98	0.81
	22	0.32	0.10	0.01			0.00	0.01	0.00	3.20			4 (3		1 85	9	0.32		0.42	0.11	0.15	0.54	0.32	01.0	0.00		0.13	0.35	0.34 >	9.26	1.57	3.85			1.34	1.15 >	0.93	0.00		9:	0.20
	oiff.	Y Y	Y Y		4	4	0.576	0.963	0.918	Y Y	M		77.		0.070		¥	¥				1.012	V V	4			0.484					0.830			1.137					0.617 <	
PL	*		:	OKH	- '			SOL			:	2					-	-	OCH O	V20	RHS			:			RJE		75X (0.03							STN		770	210	9
Test	Oate	12/04/87	03/01/88	02/06/89	987 507 20	02/01/0	02/06/89	11/29/89	12/06/90	08/11/87	05/25/88	09/20/00	02/01/02	07,110,00	00/01/80		08/11/87	05/25/88	02/07/89	03/10/89	09/01/89	11/29/89	78/60/60	05/25/88	04/10/89		09/12/89	05/31/96	07/12/90	02/28/89	03/21/89	09/27/89	91.001.00	04/15/68	02/07/89	03/10/89	12/07/89	02/07/89	03/10/89	12/07/89	02/13/90
Ship-	ment			87	2	75	87	19	61	23	25		0 0	2 0	2 %	3	S	22	48	48	52	19					35	29	29	33	23	33	,	3	48	1 00	84	48	48	89	48
	No.	2811	2811	2811	2643	7107	2812	2812	2812	2979	2070	20.00	2070	2070	2070		2980	2980	2980	2980	2980	2980	3038	3038	3038		3802	3802	3802	3966	3965	3964	020	207	4070	4070	4070	7207	4074	707	4074

< ∪ ⊢	* * * * * * *	*** ***	**: ***	* * * · · *	• • • • •	• • • • •	* * * *	
Assay Type	CPE CPE CPE CPE CPE CPE CPE CPE CPE CPE				*****			
TAI	1.10 1.40 2.35 27.24 22.13 2.46	20.16 8.99 26.83 23.73 14.47 5.15	13.40 3.41 24.56 13.42 3.69	14.99 0.00 37.38 0.35 2.00 11.23	1.90° 21.90 16.13 12.97 14.80	32.32 19.07 15.38 10.40 7.17	30.11 15.73 17.75 19.40	
15	0.40 0.00 0.00 0.00 0.00 0.00 0.00	2.24 × 4.12 × 0.79 × 1.56	2.10 × 4.39 × 2.13 × 0.00	2.61 0.00 11.20 0.00 0.00 0.00	0.07.7.0 0.07.7.0 0.09.7.4	3.87 × 1.79 × 1.80 × 0.00 × 0.00	6.54 × 3.76 × 3.26 × 3.12 ×	
A1 95	0.0000000000000000000000000000000000000	1.10 × 10.24 3.40 0.00 0.00	1.08 0.00 0.00 0.00	0.00 0.00 0.00 0.00 0.00	8 9 8 8 8 8 8 8 8 8 8 8 8 8 8 8 8 8 8 8	0.000	3.22 0.00 0.00 0.00	
70 95	0.10 1.00 0.32 0.32 2.85 0.32	10.00 × 32.00 × 100.00 100.00 100.00	10.00 × 10.00 × 32.00 × 32.00	320.00 320.00 74.60 100.00 100.00	0.00 0.00 0.00 0.00 0.00 0.00	100.00 100.00 00.00 00.00 00.00	0.10 1.00 0.89	
26 DI	0.00 0.00 0.00 0.00 0.00 0.00 0.00 0.0	9.12 9.40 9.40 0.00 0.00 0.00	9.29 0.00 0.00 0.00 0.00 0.00	0.00 0.00 0.00 0.00 0.00 0.00 0.00 0.0	× × × × × × × × × × × × × × × × × × ×	26.50 v 0.00 v 0.00 v 0.00 v	0.00	
AI 50	5.50 0.00 1.61 7.67 0.00	2.51 5.89 5.94 5.67	2.10 2.27 6.30 3.73 0.00	22.20 0.00 0.00 0.00	10.00 10.40 13.73 10.00	13.30 8.46 4.00 3.02 0.00	6.54 8.73 4.94 4.83	
10 50	0.10 0.32 0.32 0.32 0.32 0.32	10.00 × 24.60 × 32.00 × 55.20 × 22.10	10.00 × 21.60 × 12.00 7.65	320.00 320.00 33.40 100.00 28.30	10.00 × 3.75 × 2.49 × 2.49 × 8.51	92.50 50.40 71.50 76.50	0.10 × 0.21 0.31 0.26	
10 50	0.02 0.00 0.00 0.17 0.00 0.00	3.99 × 6.69 × 7.16 × 32.00 × 8.27	4.77 × 9.53 1.29 3.20 0.00	68.40 0.00 0.00 0.00 0.00 0.00	0.90 × 0.53 0.58 0.67 2.05	6.97 5.96 17.90 25.40 0.00	0.02 0.08 0.08	
AI 25	0.50 3.10 1.21 2.75 7.02 8.60	6.03 9.88 7.02 5.10 2.65	3.03 2.30 4.89 2.51	9.19 16.80 0.00 3.80 3.80	5.65 5.65 5.03 5.03 5.03 5.03	11.60 5.61 5.05 3.22 2.58	9.69 7.39 5.64 4.81	
70 25	0.10 0.32 0.32 0.32 0.32 0.32 0.32	10.00 × 15.00 × 17.10 × 25.30 × 12.90	5.66 5.82 5.54	179.00 8.36 16.80 v 100.00 19.20	3.20 - 2.17 - 1.56 - 5.06 - 5.86 -	50.10 23.10 32.00 45.60	0.10 × 0.09 v 0.21 0.17	
10.25	0.21 0.32 0.26 v 0.12 v 0.12 v	3.4.1. 8.4.4. 3.4.1. 8.9.4.	3.30 × 0.62 1.39 1.39	19.40 230.00 1.00 0.00 5.05	0.50 0.38 0.39 0.37 1.26	4.34 6.34 14.20 2.81	0.00 0.00 0.00 0.00	
Diff.	NA NA 1.210 0.827 0.809 0.700	0.745 0.926 0.680 0.697 0.956 1.015	0.697 1.015 0.706 0.956 1.081	0.945 0.655 0.859 0.744 0.764	0.744 0.744 0.615 1.006 0.899 0.705	0.764 0.615 1.006 0.965 0.734	0.810 0.830 0.934 0.725	
Test Pit Date #	05/04/88 06/14/88 02/07/89 040 03/10/89 P00 05/24/89 967 09/12/89 RJK 11/01/90 ZW	09/13/89 RLJ 05/09/90 W0Y 07/06/90 XC1 09/13/89 RLK 11/29/89 SQM 11/01/90 ZWZ	09/13/89 RLK 11/01/90 2WZ 09/13/89 RLL 11/29/89 SOH 11/01/90 2X0	07/07/69 901 08/16/69 &6Y 01/31/89 012 02/06/90 1RA 02/13/90 U01 02/13/90 1ZV	07/27/88 02/28/89 0U0 09/13/89 RLN 11/29/89 SQN 02/06/90 TRA 02/13/90 TZV	02/28/89 OU1 09/13/89 RLN 11/29/89 SQN 02/06/90 TRB 02/13/90 TZW	02/28/89 OUZ 09/27/89 RSA 12/05/89 SRV 02/13/90 TZX	
Ship- To	39 98 88 98 98 98 98 98 98 98 98 98 98 98	65 65 65 65 65 65 65 65 65 65 65 65 65 6	45 45 45 11,19	44 44 62 62 62 62 62 62 62 62 62 62 62 62 62	42 02/ 42 02/ 63 11/ 63 02/ 63 02/ 63 02/	48 42 63 63 63 63 63 63 63 63 63 63 63 63 63	48 02, 48 12, 63 02,	
No.	######################################	4277 4277 4277 4278 4278 4278	4280 4280 4281 4281 4281	4452 4452 4527 4527 4527	0654 0654 0654 0654 0654	4592 4592 4592 4592 4592	6097 6097 6097	

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Assay Type		E E	E E	FFF			ĔĔ	E E	FFF			
Ţ.	£.00.00 0.00	8.52	3.82	10.37 27.84 0.00	25.60 60.74 37.48	59.97 62.18 44.23 35.99	5.41	0.00	3.11 5.98 4.52	11.74	3.02	0.00 5.79 1.07
5	2.54 v 0.00 0.00	1.18	1.22 > 0.00 >	1.84 > 5.94 > 0.00	5.90 40.51 × 8.60 ×	23.40 > 28.79 > 13.47 > 7.35 >	3.85	3.64	0.00 1.78 0.00 v	0.00	0.00	1.37 0.00
A1 95	0000	0.0	0.00	0000	0.00 32.78 0.00	20.70 11.12 > 5.14 3.4 >	0.0	0.00	0000	00.00	0.00	0.00
TC 95	97.90 94.50 302.00	305.00	308.00	320.00 32.00 32.00	320.00 100.00 > 320.00	320.00 × 320.00 × 100.00 ×	320.00 1000.00	320.00	320.00 80.60 100.00	320.00 966.00 320.00	320.00 86.40 9.69	262.00 100.00 305.00
10 95	8888	0.00	0.00	0000	3.05 0.00	15.50 × 28.80 × 62.20 × 29.40 ×	0.00	0.00	0.00	00.00	0.00	988
AI 50	3.47	2.18	2.18	5.78 14.16 0.00	14.50 59.53 14.11	41.70 28.79 19.86 7.35	5.09	5.59	0.00 2.81 0.00	2.65	1.38 0.00 3.65	1.77
10 50	66.00 45.20 143.00 46.00	165.00 315.00	169.00 172.00	13.10 22.10 6.71	95.50 88.70 320.00	200.00 320.00 > 280.00 100.00 >	320.00	320.00	55.20 21.20 29.00	320.00 660.00 320.00	320.00 × 21.70 6.26	66.50 100.00 v 167.00
10 50	6.00 0.00 0.00	0.00	0.00	2.27 1.56 0.00	6.57	4.79 11.10 > 14.10 13.60 >	0.00 >	0.00 >	0.00 7.52 0.00	249.00	232.00 >	0.00 56.60 v
A1 25	3.72	2.11	2.15	3.25 13.55 0.00	9.73 60.33 32.99	40.50 65.23 30.31 13.94	0.00	0.00	0.70 2.78 0.90	3.30	2.10	2.03
TC 25	48.30 0.82 0.58 23.80	89.10 187.00	94.80	4.18 9.27 1.55	38.80 60.30 195.00	112.00 320.00 > 190.00 100.00 >	282.00 756.00	320.00	19.10 13.40 19.50	309.00 490.00	320.00 12.80 0.32	39.60 77.30 92.70
10 25	13.00 0.00 0.00	42.30	00.00	1.29 0.68 0.00	3.99 1.00 5.91	2.77 4.91 > 6.27 7.17 >	0.00	0.00 100.00	27.30 4.82 21.70	148.00	152.00 5.21 0.62	38.10
Diff.	0.742 0.655 0.398 0.873	1.017	0.954	1.129 0.490 1.081	1.080 0.539 0.916	0.598 0.615 0.975 1.032	0.960	0.873	0.887 1.087 0.842	1.017	1.085	0.955 1.029 0.976
¥	9 802 9 864 0 UG3	8 02a 0 003	8 02R 0 U03	9 OF2 9 RLP 0 2X0	9 00% 9 810	9 80ET 9 SOR 0 1PG	9 SAS	9 P00	9 POR 9 STU 9 SXR		9 S1X 9 OSF 9 S1Y	9 OSK 9 S12 0 TFU
Test	07/07/89 08/16/89 03/08/90 05/03/90	11/08/88 02/13/90	11/08/88 02/13/90	01/24/89 09/13/89 11/01/90	01/24/89 09/13/89 11/29/89	02/21/89 09/13/89 11/29/89 12/06/90	01/16/89	03/13/89 12/05/89	03/13/89 10/10/89 12/14/89	03/13/89 11/29/89 03/13/89	10/10/89 02/27/89 10/10/89	02/27/89 10/10/89 01/18/90
Ship- ment	3333	22	22	333	332	87 5 87	97	51	222	2 2 2	S 87	84 84
AVS.	\$55, \$55, \$55, \$55, \$55, \$55, \$55, \$55,	8927	6927	4785 4785 4785	4796 4796 4796	4855 4855 4855 4855	12871	8267	7867 7867 7867	7867 7885 7887	\$032 \$032	5053 5053 5053

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Assay Type				H	FF	FE	FF	HIN	HHH		FFFF		
IĀĪ	8.39 1.37 3.81 5.65	3.42	20.58 22.70 19.41	30.37 23.50 20.59	1.47	0.00	10.78	18.89 13.48 7.97	5.44 0.92 0.00	6.26 0.00	11.51 13.21 7.68 4.02	17.62 16.05 0.09 0.23	
15		0.00	3.22	0.16 × 4.39 4.65 ×	0.00 > 2.76	3.09	1.99 > 2.52 >	2.29	0.00	1.14 0.00 0.00 0.00	1.85 \$ 1.02	0.00	
A1 95	98888	0.00	3.32	3.39	0.00	8.08	0.0	0.00	888	9000	0.000	0.000	
75 %	320.00 320.00 1000.00 1000.00	320.00 320.00 977.00	309.00 309.00 320.00 >	320.00 × 1000.00 × 1000.00	309.00	7.91	91.50	1000.00 × 1000.00 320.00	32.00 32.00 100.00	320.00 1000.00 3200.00	25.10 32.00 100.00 95.40	320.00 1000.00 981.00 320.00	
10 %	98888	0.00	93.00 91.50 288.00 >	94.30 × 293.00 × 0.00 ×	0.00	0.00	0.0	914.00 v 0.00 v 0.00 v	0.00	0.00	0.000	0.000	
AI 50	1.03 0.00 0.00 0.00	0.00	4.36 5.11 2.88	5.76 6.33 4.65	3.74	0.00	3.13	3.17	0.00	0.00	4.04 0.00 1.80	0.00 0.00 0.00	
10 50	320.00 320.00 1000.00 1000.00	320.00 × 215.00 152.00	210.00 210.00 320.00 >	320.00 × 827.00 1000.00 ×	36.20	0.73	2.66	1000.00 × 715.00 320.00 ×	32.00 × 21.60 6.94	320.00 × 915.00 2210.00	5.75 32.00 71.30 54.10	320.00 767.00 550.00 320.00	
10 50	310.00 0.00 0.00 0.00 0.00 0.00	207.00 \$ 0.00 0.00	48.20 41.10 111.00 >	55.60 × 131.00 × 215.00 ×	0.00	0.00	0.85	405.00 > 226.00 > 237.00 >	25.80 v 0.00 0.00	282.00 \$ 0.00 0.00	0.00 × 27.90 × 30.10	221.00 0.00 0.00	
AI 25	2.32 0.00 1.77	0.00 0.00 0.00	4.64 7.22 6.14	0.22 7.95	0.00	0.00	3.57	4.10	0.00	0.00	2.63 3.77 2.24	11.30 4.57 1.14 1.13	
TC 25	> 320.00 > 320.00 > 1000.0 > 1000.0	88.00 139.00 47.70	155.00 155.00 > 320.00 >	8.95 574.00 > 1000.0 >	11.80	< 1.00 0.53	1.69	> 1000.0 > 518.00 283.00	28.30 15.80 5.07	> 320.00 > 561.00 1430.0	2.63 × 44.80 × 30.60	166.00 544.00 306.00 282.00	
10 25	138.00 0.00 566.00 234.00	134.00 0.00 0.00	33.40 21.50 52.10	41.40 72.20 147.00 3	36.70	0.00	77.0	204.00 126.00 103.00	14.30 0.00 0.00	0.00		14.70 119.00 269.00 249.00	
Diff.	0.881 1.029 0.993 0.857	0.724 0.935 0.895	0.623	0.830 0.542 0.561	1.007	0.808	0.635	0.849 0.872 0.835	0.864 0.935 0.667	0.927	0.798 0.960 0.874 0.875	0.788 0.947 0.942 0.887	
£ *	9 OSL 9 S1Z 9 SR9 9 T15 0 TVI	9 0TZ 9 RS7 9 RUM	9 GHH 9 R21 0 ZX1	9 9PM 9 R62 0 V01	9 oP0 9 R70	0 x2v	0 V25	52V 0	9XX 0	0 V27 0 XCS 0 ZZE	9 PKC 9 SXM 0 VZ9 0 XKE	9 PKD 9 T19 0 VRS	
Test	02/27/89 10/10/89 12/05/89 12/20/89	02/28/89 09/27/89 10/03/89	06/21/89 08/08/89 11/01/90	07/11/89 08/16/89 04/26/90	07/11/89 08/16/89	05/08/90	05/08/90 07/18/90	12/12/89 12/14/89 05/08/90	12/14/89 05/08/90 07/18/90	05/08/90 07/06/90 11/07/90	04/18/89 12/14/89 05/08/90 07/18/90	04/18/89 12/20/89 05/01/90 05/08/90	
Ship-	2 2 2 2 8 8 8 8 8 8 8 8 8 8 8 8 8 8 8 8	999	222	57 57 65	57	33	88	223	288	333	8888	8888	
AVS	5058 5058 5058 5058 5058	502 502 502	\$121 \$121 \$121	5138 5138 5138	5142	2075	5483	7875 7875 7875	5485 5485 5485	6875 6875 6875	222 223 223 233 233 233	\$497 \$497 \$497	

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Assay Type	EEE			E E	# H H		EFFF		TIM	ĒĒ	FF	F F
IĀ	17.26	21.27 4.83 1.91 8.75	21.73	1.37	18.67 23.28 9.06	8.88 8.88 8.69 11.88	0.43 17.12 0.06 0.00	8000	16.14 18.09 0.00	8.02	11.52 0.19	3.51
18	2.98 0.00 v	3.04 0.00 0.00 v	2.85 4.4.4		4.10 × 0.00	> 313.00 > 313.00 > 313.00 > 313.00 > 5.87 >	0.00 0.00 0.00 0.00	0.00	3.58 > 1.47 > 0.00	1.78 > 0.71 >	0.00	5.23
A1 %	98.0	0000	88 88	0.0	32.40 29.48 0.00	× 313.00 × 313.00 0.00 0.00	9888	0.00	0.00	0.00	9.0	9.8
70 %	295.00 309.00 302.00	163.00 309.00 300.00 297.00	320.00 973.00 320.00	320.00	100.00 94.30 32.00	100000	1000.00 3200.00 1000.00 3200.00	320.00 320.00 241.00	320.00 32.00 32.00	30.70	31.30	320.00 100.00
1C %	90.0	90000	98 98	0.0	3.20	< 320.00 > 0.32 > 0.00 > 0.00 >	00000	0.00	0.00	0.00	0.00	0.00
AI 50	0.00	5.20 0.00 0.00	3.85	2.00	6.72 5.49 0.00	313.00 < 313.00 < 313.00 < 5.87	2.23	1.23	5.38	2.70	0.00	21.47
TC 50	154.00 210.00 140.00	58.40 74.40 82.70 97.40	320.00 663.00 320.00	320.00 587.00	9.28 7.35 6.00	100000	1000.00 3200.00 806.00 1690.00	282.00 32.00 66.00	9.62 32.00 3	19.10	3.20	75.30 13.70
10 50	8.00	0.00	204.00 >	0.00 >	1.38	320.00 > 0.32 > 320.00 > 17.00 >	0.00 × 0.00 × 0.00 0.00	230.00	21.70 >	3.20	2.12	3.51
8	9.68 2.18 1.29		10.51		2.5. 2.6. 2.6.	313.00 < 313.00 < 11.40	0.00 7.00 7.00 7.00	0.00	6.41	2.65	3.01	18.34
70 25	76.10 155.00 71.20	34.20 49.70 53.90 64.70	320.00 × 492.00 × 294.00		5.66 5.27 2.90	100000	627.00 3200.0 228.00 622.00	10.00 < 1.00 42.40	6.41 32.00 16.50	12.60	4.25	18.30 ×
10 25	7.87 71.00 55.20	5.05 0.00 0.00	156.00 × 46.80 × 105.00	224.00 >	1.000.551.10	< 320.00 > 0.32 > < 320.00 > < 320.00 > 8.77 >	0.00 521.00 0.00 850.00	0.00	< 1.00 4.91 × 0.00	¥.5.	1.41	1.00
Diff.	0.703	0.639	0.985	0.953	0.970	0.486 0.486 0.655 0.655	0.806 0.769 0.929 0.810	0.937	0.800 0.605 0.810	0.787	0.787	0.940
¥ 5 6	8/89 PKG 3/90 UOS 8/90 VZC	8/89 PKH 8/90 1FU 0/90 W1J 1/90 ZX2	4/89 90K 9/89 R3P 4/89 90M		4/89 000 9/89 R3R 1/90 2X2	3/89 PY8 3/89 PY8 1/89 056 1/89 056	5/89 SRR 8/90 TWJ 1/90 WGU 1/90 ZX3	6/89 QMC 6/89 R75 1/90 VRU	8/89 QT4 2/89 R8V 1/90 2X3	1/89 SAC 0/90 TN1	10/31/89 SAC 11/30/90 TN2	11/07/89 SES 11/01/90 2X4
. Test Date	04/18/89 02/13/90 05/08/90	04/18/89 01/18/90 05/10/90 11/01/90	06/14/89 08/09/89 06/14/89	06/14/89	06/14/89 08/09/89 11/01/90	05/03/89 05/03/89 05/11/89 05/11/89	12/05/89 SN 02/08/90 05/31/90 67 11/01/90	07/06/89 08/16/89 05/01/90	07/18/89 08/22/89 11/01/90	10/31/89 01/30/90	10/31/89 01/30/90	11/07/89
Ship-	233	22.32	22 22	2 %	888	2222	62 CABSN 67 62/67	57 57 65	57 57 57	2 2	2 6	2.2
AVS.	5503 5503 5503	5515 5515 5515 5515	5532 5532 5532 5535	5538	5539 5539 5539	5580 5580 5580	5601 5601 5601	5643	5695 5695 5695	5905	5906 5906	5997 5997

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Assay Type			T T T	TH TH	T I I	THE	TH	TIM	TIM	T I I	H	T I I	FFFF		FF	
IĀ	14.27	11.80 29.00 23.98	10.08	13.49	7.56	16.40	14.73	12.46	10.75	25.51 29.31	23.56	12.08	0.00	3.42 5.16 0.49	0.00	
Is	4.41 v 18.94 v 2.77	3.80 \	1.83	0.00	1.46 \ 0.00 \	3.08	2.74	2.57	1.63 > 2.74 >	3.64	3.00 >	1.3° 2.3°	0.00	0.00	0.00	
A1 95	888	1.07 × 3.31 3.17	3.15	3.39	0.00	3.39	0.00	0.0	1.06 >	3.40	1.12 >	1.06 >	8808	0.00	0.00	
TC 95	320.00 32.00 32.00	320.00 × 966.00 966.00	320.00 > 952.00	320.00 > 796.00	320.00	320.00 > 309.00	%.60 %.20	309.00	320.00 >	320.00 >	320.00 >	320.00 >	1000.00 3200.00 3200.00 3200.00	96.60 96.10 96.10	262.00	
55 56	0.00	300.00 × 292.00 × 304.00	302.00 >	94.50 >	0.00	94.50 >	30.20	0.00	301.00 >	290.00 > 294.00 > 1	287.00 >	302.00 >	0.00	0.00	0.00	
AI 50	20.56 18.94 8.15	1.86 5.12 4.61	2.91	3.84	1.46	0.00	3.69	3.48	1.83	5.28	3.00	5.5 5.5	0.00	0.00 2.17 0.00	0.00	
10 50	22.20 32.00 > 22.80	320.00 > 660.00 657.00	320.00 > 521.00	217.00	320.00 > 630.00	249.00	66.00	210.00	320.00 >	320.00 >	320.00 >	320.00 >	1000.00 3200.00 > 2680.00 3200.00	66.00 61.40 60.90	158.00	
10 50	1.08 1.69 × 2.80	170.00 × 129.00 142.00	179.00 \$	56.60	219.00 > 0.00	56.60	0.00	60.30	175.00 \$	121.00 >	107.00 > 87.10 >	179.00 > 171.00	2670.00 × 0.00 × 0.00 × 0.00 × 0.00	0.00 28.30 0.00	32.00	
81 R	5.83	2.58 10.73 6.57	2.39	3.73	2.16	6.10	3.66	3.53	3.66	8.27	8.71	2.39	0.00	1.93 2.76 0.00	0.00	
TC 25	4.77 × 32.00 7.73	320.00 × 490.00 × 485.00	320.00 > 290.00	159.00	320.00 > 446.00	174.00 155.00	49.00	155.00	320.00 >	320.00 >	320.00 >	320.00 >	755.00 3200.0 > 1230.0 1920.0	49.00 42.00 41.40	19.70	
10 25	1.00	124.00 × 45.70 73.80	134.00 ×	42.50	148.00 > 239.00	42.50	13.40	0.00	129.00 \$	38.70 v 58.80	54.30 >	134.00 > 125.00	0.00 1080.0 > 0.00 2780.0	25.40 15.20 0.00	0.00	
Diff.	0.919 < 0.849 0.956	1.066 0.880 0.916	1.066	0.865	0.830	0.853	0.853	0.776	0.825	0.998	1.082	0.971	0.916 0.827 0.894 0.989	0.875 1.018 0.989	0.948	
=	9 SES 0 TOR 0 TWA	5 × × ×	0 TA0	0 TA0	0 TAR	20Z 0	20Z 0	0 TAT 0 2V0	0 ZV0	1CH 0 2X4	0 TCK	0 TCN	SRS TWK 2X6	100 CO	0 VIR	
Test	11/07/89 SES 02/01/90 TOR 02/08/90 TWA	01/11/90 10/31/90 12/06/90	01/11/90	01/11/90	01/11/90	01/11/90	01/11/90	01/11/90	01/11/90	01/16/90	01/16/90	01/16/90	12/05/89 02/08/90 05/31/90 11/01/90	02/27/90 03/20/90 11/01/90	05/10/90	
Ship-	222	333	33	33	29	29	23	33	33	2 23	2 29	33	62 GABSH 67 67	222	33	
AVS S	9665 9665	818 818 818	6196	6199	6202 6202	6203	6204	9029	6207	6214 6214	6219 6219	6225	6234 6234 6234 6234	6369 6369 6369	£13	

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Table 12 (Cont'd)

∀ ∪ ⊢	. +	+ •	+ +	+ +	+ +	+ +	+ +		+ + +	+ + +	+ + +	+ 1	+ +	+ +	+ +	+ +	
Assay																	
AS Y	EE	FF	FF	H	FF	FF	H	ĒĒ	EEE	EEE	EEE	FF	EE	FF	FF	EE	
TA.	0.00	07.0	35.50 32.56	20.71	16.70 10.70	16.01	0.52	0.00	0.87 7.55 1.19	5.2 5.72 7.29	8.05 8.08 10.98	00.0	14.47	0.55	3.39	8.72	
ıs	0.00	1.12 > 0.00 >	10.40	0.00 >	1.86 >	0.00	0.00	0.00	0.53	0.00 \$	0.00	0.00	2.87 > 0.00	0.30	1.58	1.02 \$	
AI 95	0.00	0.00	33.91	0.00	1.06	3.22	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00 \	
70 95	96.30 251.00	1000.00	320.00 >	320.00	320.00 >	320.00 973.00	320.00	320.00 1000.00	309.00 307.00 320.00	100.00 308.00 100.00	100.00 100.00 100.00	32.00	320.00	320.00	320.00	1000.00	
10 %	0.00	0.00	9.44 v	0.00	301.00 > 316.00 >	302.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	3020.0 > 3	
AI 50	3.19	0.00	41.15	2.14	1.86	3.71	0.00	0.00	1.51 2.29 0.00	0.00	3.70	1.84	4.19	1.02	2.81	1.02	
10 50	40.70	1000.00 > 3200.00	163.00	320.00 205.00	320.00 >	320.00 663 .00	320.00 >	320.00	144.00 191.00 192.00	44.00 84.30 65.30	30.00 94.10 100.00	18.40	86.80 320.00	320.00 >	73.10	1000.00 > 3200.00 >	
10 50	0.00	814.00 v	3.98	0.00 v 95.90	172.00 >	0.00 \$	288.00 > 0.00	0.00 > 749.00 >	95.20 83.30 0.00	0.00	8.11 0.00 0.00	0.00	20.70	313.00 > 0.00	26.00 32.00	978.00 > 1790.00 >	
AI 25	6.00	1.78	16.83	21.72	2.53	3.68	1.02	0.00	2.67	1.43 2.32 2.69	1.94	1.93	4.42	0.53	2.65	1.79	
TC 25	13.10	909.00	35.10	320.00 > 148.00	320.00 >	220.00	173.00	320.00	\$0.90 126.00 88.70	21.80 40.70 32.00	8.79 29.00 25.70	8.37	59.40 320.00	93.20	41.00	1000.0 >	
S2 21	0.00	510.00	2.08	14.70 >	127.00 >	235.00	170.00	490.00	\$1.20 47.10 61.10	15.30 17.60 11.90	4.54 14.90 5.31	4.33	13.40	177.00	15.50	559.00 >	
Diff.	0.888	0.878	0.728	1.001	0.765	0.779	0.899	1.041	1.007 0.747 0.902	0.976 0.656 0.811	0.767	0.776	0.753	0.855	1.049	0.870	
Test Pit Date #	05/15/90 W3Y 07/11/90 XEE	04/19/90 VFK 06/27/90 X4G	06/21/90 WZC 07/12/90 XGA	07/24/90 XQJ 10/24/90 ZNT	07/26/90 XTA 10/24/90 ZNV	06/21/90 LYJ 08/09/90 XZT	08/02/90 XYB 09/27/90 243	08/30/90 YCR 10/03/90 272	08/31/90 YEX 10/03/90 278 10/25/90 296	08/31/90 YEY 10/03/90 279 10/25/90 297	08/31/90 YEZ 10/03/90 27A 10/25/90 298	08/15/90 Y3S 09/13/90 YT0	08/15/90 Y3T 09/13/90 YT1	08/23/90 Y7V 09/20/90 YV4	08/30/90 YC0 10/03/90 270	09/20/90 YXG 10/25/90 ZQB	
Ship-	33	33	79	\$ \$	69	33	66	22	222	222	222	22	22	22	22	R R	
AVS S	£13	7199	7219	6942	9769	9869	7045	7051	7083 7083 7083	7085 7085 7085	7087 7087 7087	7319	7321	25 25 25	7433	7445	

				degree of surements.		d by using		ion values	9-320).		
∢ ∪⊢	• •	• •	• •	ant the		lculate	×i	reduct	ale is		
Assay Type	EE	FE	TIM	into accourate with		or 95% ca	50% or 95	50% or 95%	EXIMM SC		CPE).
¥	1.85	1.10	4.13	takes e accu		5x, 503	y 25%,	h 25x,	, the		tion ir
ï	0.00	0.00 >	0.00 > 4.13 2.65 > 15.09	vity that TAI is mor		I CPE by 2	fability b	ulated wit	one, #9/ml	100%).	25% reduc
8	88	9.0	88	irel active to TAI.	ities.	bited vira	v lles bes	xand, calca	10 dilutio	cale of 0	st (f.e.
56 21	0.00 > 320.00	0.00 > 320.00	0.00 > 320.00	tive antiv	ical dens	that inhi	that redu	the compo	e-half-log	d upon a s	nactive te
اد %	9.0	9.0	0.0	of selections of compounds of the compound of the compounds of the compound of the compounds of the compound of the compou	antrol opt	on (kg/ml)	(m/8#)	effect of	upon 6 on	ves (base	otes an i
A1 50	0.00	0.00	3.95	asurement by the to	e virus co	ncentratí	centration	icellular	pesed) 09	iviral cur	A "-" der
10 50	98.60 158.00	3200.00 >	320.00 938.00	R is a me y produced	of and th	he drug co	drug con	al and ant .95).	by the IC	d the ant	in CPE.
10 50	90.20	0.00 > 320.00 2220.00 > 3200.00 >	0.00 > 320.00 238.00 938.00	This value is a virus rating (VR) rather than a TAI. The VR is a measurement of selective antiviral activity that takes into account the degree of inhibition of virus-induced CPE and the degree of cytotoxicity produced by the test compound similar to TAI. TAI is more accurate with MTI measurements.	the cell control and the virus control optical densities.	50% and 95% = The drug concentration (μ g/ml) that inhibited viral CPE by 25% , 50% or 95% calculated by using refitting.	and $95x$ = The drug concentration ($\mu g/ml$) that reduced cell viability by $25x$, $50x$ or $95x$.	of the antiviral and anticellular effect of the compound, calculated with 25 %, 50 % or 95 % reduction values by the $1C_{25,50,95}$).	Selectivity index = A ratio calculated by dividing the IC ₂₅ by the IC ₅₀ (besed upon 6 one-half-log ₁₀ dilutions, £9/ml, the maximum scale is 0-320).	Total Antiviral Index = The area between the cytotoxicity and the antiviral curves (based upon a scale of 0-100%).	>25% reduction in CPE. A *** denotes an inactive test (i.e. <25% reduction in CPE).
AI 25	0.00	0.00	0.00	than a T	ce in the	25%, 50% of curve fi		ratio of t	, dividing	the cytot	
TC 25	22.80	0.00 > 320.00	0.00 > 320.00	This value is a virus rating (VR) rather than inhibition of virus-induced CPE and the degree	The differential is the difference in	(Viral) inhibitory concentration 25%, 5 a regression analysis for semilog curv	(Cell) toxicity concentration 25%. 50%	Antiviral Index = A single point ratio (calculated by dividing the TC _{25,50,95}	culated by	a between	Activity = A "+" denotes a test that produced
10 25	0.00 53.70	0.00 >	0.00 ×	rating (V	al is the	tory conc	y concent	x = A sing	ratio cal	= The are	es a test
	0.701	0.625	0.825	virus irus-in	ferenti	inhibi	toxicit	ated by	ex = A	Index	" denot
Plt biff.	YXM ZUS		2E2 229	n of v	The di	Viral	(Ceff)	Intivi	ty Ind	iviral	- V =
Test	09/20/90 YXH 10/31/90 ZUS	10/11/90 ZEZ 11/07/90 ZZ9	10/11/90 ZEZ 11/07/90 ZZ9	This valuabilities					Selectivi	Total Ant	Activity
Ship-	RR	2K	2K								
No.	7461	200	£ £		7	\$6.0	96'0	56,05	W	W	10
					IFRHTL	1625,50,95	1625,50,95	1 25,50,95	=	¥	CT

4.1.5 Venezuelan Equine Encephalomyelitis Virus (VE):

The number of single drug tests carried out against VE during this contract period is summarized in yearly increments in Figure 24. During this five-year period two main in vitro antiviral assay protocols were implemented:

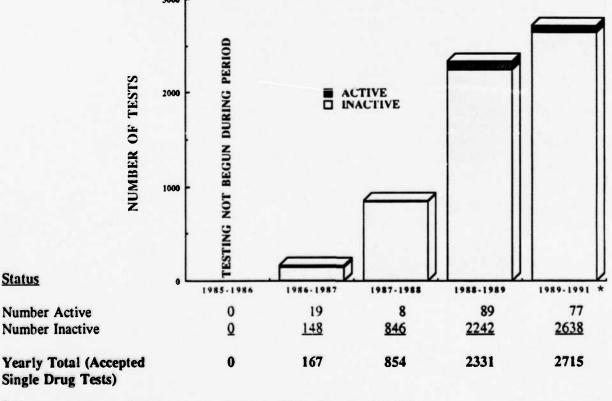
- A standard CPE inhibition assay by virus rating (VR) (Annual Report, December 15, 1. 1988, Section 3.2.4).
- 2. Since November, 1988, MTT based-antiviral assay format.

A total of 7319 tests were performed during this contract period using both assay types. Routine testing was changed to the MTT-assay format to improve the efficiency and quality of the primary screening program in addition to being more cost-effective. Selenazofurin (AVS-0253) was tested in each standard virus rating (VR) CPE-inhibition assay as a positive control compound. Results of these positive controls (VR tests) were used as a guideline to assess the quality of each assay.

After the testing was converted to the MTT-assay format, we performed a total of 406 control compound assays with Selenazofurin during the last 26 months of the contract period. During this time 684 tests were internal (+ + +) virus load, cell load, and other quality control tests. Four hundred thirtyeight (438) tests were considered unsatisfactory based on the criteria of the quality controls set during this reporting period. The rest, totaling 4756 were actual single drug MTT-assays. The total number of MTT-assays (6284) tested during the last two years represents a 224% increase (improvement) in the total testing output as compared to the total of 1937 tests performed during the first 3 years of this contract.

Out of the 6067 accepted single drug tests, 193 compounds demonstrated antiviral activity at greater than 50% reduction levels. This represents around 3.0% of the tested compounds having in vitro antiviral activity against PT-virus. The remainder, 6058 compounds (97%), were considered inactive with both assay protocols (Figure 24).

IN VITRO PRIMARY SCREEN: NUMBER OF COMPOUNDS FOUND ACTIVE AGAINST VENEZUELAN EQUINE ENCEPHALOMYELITIS VIRUS DURING THE CONTRACT PERIOD



Five-Year Totals

193

5874

6067

Status

Number Active Number Inactive

Represents 14-month period (November 15, 1989 - January 31, 1991)

- 4.1.5.1 <u>VE-Quality Controls:</u> Two positive control compounds (Schenazofurin and 2-Thio-6 Azauridine) were used in the daily assay sets as antiviral activity quality controls. The antiviral performance of the unknown compounds was compared to that of the positive control compounds. Compounds with equal to of better antiviral potency are considered active and are worthy of further in vitro profile studies and in vivo testing.
- 4.1.5.1.1 Antiviral Activity of Selenazofurin vs VE Virus: A summary of the antiviral and cytotoxicity performance of the primary control compound, AVS-0253 (Selenazofurin) is presented in Figure 25-A for 220 tests performed during November, 1989 through January, 1991.

Control Compound-Antiviral Performance: Selenazofurin (AVS-0253) has been the sole control compound against VE in these MTT-assay screens. The mean and median antiviral inhibition and cytotoxicity patterns of the positive control drug (Selenazofurin) are illustrated in Figure 25-A.

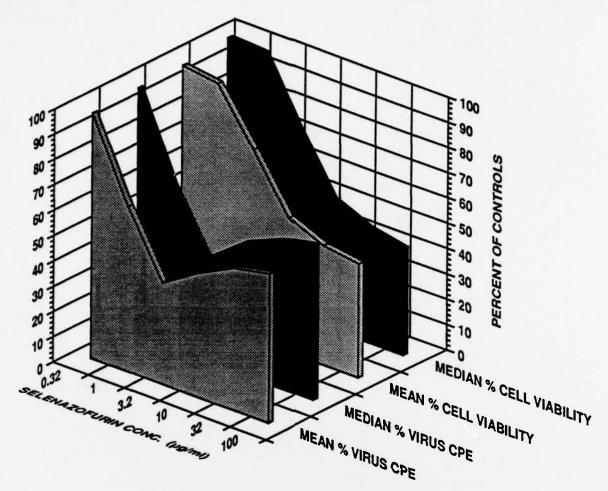
The 220 control tests performed with Selenazofurin gave a mean Total Antiviral Index (TAI) of 17.10% (SD \pm 14.12) and the median value was 13.13%. The TAI measures the overall antiviral effectiveness of the compound and it ranged from ~ 0.00 - 64.83% during this period. The mean Selectivity Index (SI) was 4.52 (SD \pm 9.59) and the median SI value was 1.75, indicating poor antiviral selectivity for Selenazofurin and it ranged from ~ 0 - 88.38% during this period. However, the closeness of the mean and median values indicate that the present execution of the SOP is consistent and repeatable.

The mean Antiviral Index 25% (AI₂₅) value was 9.25 (SD \pm 16.22). The median AI₂₅ value was 4.42 (range 0.3 - 137.38). The mean Antiviral Index 50% (AI₅₆) was 28.03 (SD \pm 41.56) with a median of 7.92 (range 0 - 254.46). The mean Antiviral Index 95% (AI₅₆) was 1.7 (SD \pm 10.12) with a median value of 0 (range 0 - 109.79). This indicates that the control compound, Selenazofurin, does not consistently reach 95% antiviral reduction levels.

The mean Antiviral Inhibitory Concentration 25% (IC₂₅) was 0.87 μ g/ml (SD \pm 0.53). The median IC₂₅ value was 0.71 μ g/ml (range 0.22 - 5.52 μ g/ml). The mean Antiviral Inhibitory Concentration 50% (IC₅₀) was 1.44 μ g/ml (SD \pm 1.28). The median IC₅₀ value was 1.40 μ g/ml (range = 0 - 10.00 μ g/ml). The mean Antiviral Inhibitory Concentration 95% (IC₉₅) could not be attained with Selenazofurin versus Venezuelan Equine Encephalomyelitis Virus. This discrepancy indicates that the control compound Selenazofurin does not consistently reach 50 or 95% reduction levels.

The average maximum antiviral inhibitory level of 220 Selenazofurin tests (Figure 25-A) was reached at 3.2 μ g/ml of the compound with 60% antiviral effect. Further increase of the drug concentration does not improve its antiviral activity. Maximum antiviral effect (~60%) was found with a simultaneous ~20 - 25% cytotoxic suppression. Above the 10 μ g/ml concentration the antiviral protection levels off to ~40% reduction level at 100 μ g/ml while simultaneously the Selenazofurin becomes maximally toxic (~60%).

SELENAZOFURIN - VS -- VE VIRUS



CONCENTRATION (µg/ml)

		%	Viral	CPE					% Cel	l Viabil	ity	
Conc.(µg/ml)	0.32	1	3.2	10	32	100	0.32	1.	3.2	10	32	100
Mean	97	65	40	49	56	59	98	96	75	53	47	45
Median	98	65	41	52	58	61	100	99	74	51	45	42
Std. Dev.	0.05	0.16	0.16	0.13	0.11	0.10	0.04	0.07	0.14	0.13	0.11	0.11

Figure 25-A
Average Antiviral and Cytotoxicity Values for 220 Positive Control Compound Tests

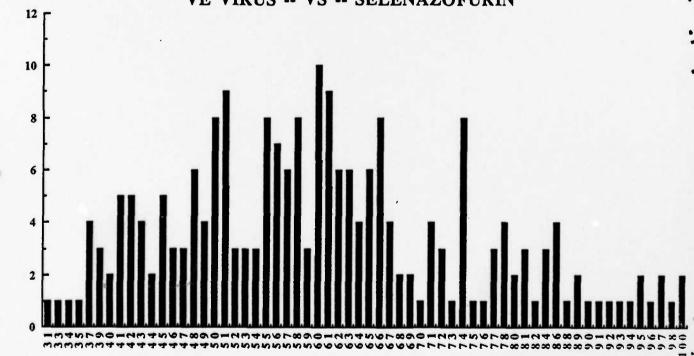
4.1.5.1.2 <u>Maximum Antiviral Effect of Selenazofurin vs VE Virus (VE)</u>: Since the metabolic activity of the cells was an unknown function during the testing period, it was monitored indirectly by measuring the maximum antiviral effect of the control compound Selenazofurin. This demonstrated the amount of infectious virus that was produced by the cells (Maximum Percent CPE).

A bar graph scatter plot (Figure 26-A) depicts the distribution of the maximum antiviral reduction values of all 220 control compound assays for Selenazofurin. The results indicate that the average maximum antiviral reduction obtained with the present SOP is around 61.0% (SD \pm 15.13) reduction levels. The maximum reduction levels vary from 31 - 100% but remain quite consistently around the median of 60%. The assay control values give a relatively broad, bell-shaped distribution curve. This indicates quite a consistent day-to-day performance of the control compound in the VE-MTT assay.

During this period the positive control compound performance criteria for Selenazofurin versus the VE virus was set at 25% reduction level. All assays in which Selenazofurin did not meet this accepted quality control level (\geq 25%) were rejected (i.e., 161 unsatisfactory tests).

Since Selenazofurin is only marginally active against VE virus, better quality control compounds are needed. However, regardless of the poor performance of the VE quality control drug Selenazofurin, around 323 different compounds have equal or better antiviral activity against VE virus than AVS-0253. Some of these could certainly be used as a better *in vitro* antiviral control compound in this large-scale antiviral screening program.





PERCENT CPE REDUCTION

Figure 26-A

Maximum Antiviral CPE Reduction (%).

Summary of 220 Control Tests.

147

4.1.5.1.3 Cellular Cytotoxicity of Selenazofurin vs VE Virus:

VE-Control Compound-Cytotoxicity Performance: The 220 cytotoxicity values of the positive control compound Selenazofurin are also very consistent. The mean cell Toxic Concentration 25% (TC₂₅) was 6.03 μ g/ml (SD \pm 9.44) and the median was 3.11 μ g/ml (range of <0.32 - 90.3 μ g/ml). The mean cell Toxic Concentration 50% (TC₅₀) value was 36.53 μ g/ml (SD \pm 39.24) and the median was 13.3 μ g/ml (range of 2.15 - > 100 μ g/ml). The mean cell Toxic Concentration 95% (TC₉₅) value cannot be attained with Selenazofurin versus Venezuelan Equine Encephalomyelitis Virus.

As can be seen from Figure 25-A, the toxicity starts to become measurable above the concentration of 1.0 μ g/ml and the maximum toxicity has not been reached at 100 μ g/ml.

When the cytotoxicity reaches around 50% (10 μ g/ml), the control compound (Selenazofurin) has reached close to its maximum toxicity. After 3.2 μ g/ml the antiviral protection of Selenazofurin starts to decrease down to ~40%. Selenazofurin becomes maximally toxic between 10 - 100 μ g/ml. The highest Selenazofurin concentration tested in these assays was 100 μ g/ml.

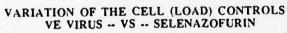
Selenazofurin has a definite cytotoxic suppression on cellular metabolism and growth. The TC_{25} and TC_{50} toxicity can be achieved with relative consistency at 100 μ g/ml. A peculiar feature of the control compound is that the cytotoxicity levels off after reaching its maximum cytotoxic value ($\pm 50\%$) at 10 μ g/ml and continues at that stationary level at increasing drug concentrations.

4.1.5.1.4 <u>VE-Assay Plate Ouality Controls: Cell Load and Virus Load Parameters (Selenazofurin):</u> The MTT assay is fundamentally dependent upon the quality of the assay plates. Our large-scale antiviral testing is dependent upon the uniformity of the test plates produced for the daily assays. Equal numbers of cell load and virus load as well as the consistent performance of the reagents used daily was monitored. A sample of the plate variation control for the period of November, 1989 through January, 1991, is presented in Figures 27-A, 28-A and 29-A.

VE-Control Compound-Cell Load Performance: A bar graph scatter plot of the mean cell control (O.D. reading) of 220 control assays is plotted in Figure 27-A. The results indicate that the cell O.D. readings reached a mean 1.160 (SD \pm 0.150) with a median of 1.150 (range of 0.845 - 1.579). This indicates that a uniform and equal number (18,000 cells/well) of cells are being loaded into every well in the 96-well plate during the day-to-day operation. The cells reduced MTT to formazan giving maximum blue color uniformly and consistently.

<u>VE-Control Compound-Virus Load Performance</u>: A bar graph scatter plot of the mean virus load O.D. readings of the 220 control assays is presented in Figure 28-A. The results indicate that the average virus load O.D. reading is 0.038 (SD \pm 0.037) with a median of 0.029 (range of 0.012 - 0.229). This demonstrates that a good cell destruction is taking place and a uniform load of virus (32 TCID₅₀) is administered on the cell monolayer with very consistent viral CPE results.

<u>VE-Control Compound-Assay Differential Performance:</u> A bar graph scatter plot of the mean O.D. differential values of the 220 control assays is provided in Figure 29-A. The results indicate that the average differential O.D. reading is 1.122 (SD \pm 0.149) with a median of 1.116 (range 0.691 - 1.520). The single bell-shaped curve is reasonably sharp and uniform. This reflects that the assays are executed consistently and are repeatable during day-to-day operation with close to 82% measurement accuracy.



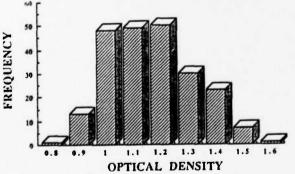


Figure 27-A

VARIATION OF THE VIRUS (LOAD) CONTROLS VE VIRUS -- VS -- SELENAZOFURIN

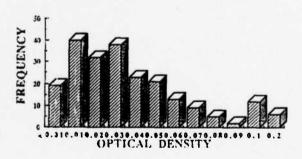


Figure 28-A

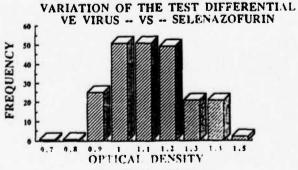


Figure 29-A

4.1.5.1.1 Antiviral Activity of AVS-6724 (2-Thio-6-Azauridine) vs VE Virus: A summary of the antiviral and cytotoxicity performance of the second control compound, AVS-6724 (2-Thio-6-Azauridine) is presented in Figure 25-B for 42 tests performed during November, 1989 through January, 1991.

Second Control Compound-Antiviral Performance: 2-Thio-6-Azauridine (AVS-6724) has been tested as a possible second control compound against VE in these MTT-assay screens. The mean and median antiviral inhibition and cytotoxicity patterns of this second positive control drug are illustrated in Figure 25-B.

The 42 control tests performed with 2-Thio-6-Azauridine gave a mean Total Antiviral Index (TAI) of 2.50% (SD \pm 6.40) and the median value was 0.17%. The TAI measures the overall antiviral effectiveness of the compound and it ranged from \sim 0.00 - 26.84% during this period. The mean Selectivity Index (SI) was 0.51 (SD \pm 1.60) and the median SI value was 0, indicating poor antiviral selectivity for 2-Thio-6-Azauridine and it ranged from \sim 0 - 7.43 during this period. However, the closeness of the mean and median values indicate that the present execution of the SOP is consistent and repeatable.

The mean Antiviral Index 25% (AI₂₅) value was 1.10 (SD \pm 2.50). The median AI₂₅ value was 0.23 (range 0 - 12.49). The mean Antiviral Index 50% (AI₅₀) was 5.70 (SD \pm 20.10) with a median of 0 (range 0 - 110.48). The mean Antiviral Index 95% (AI₉₅) was 2.47 (SD \pm 16.00) with a median value of 0 (range 0 - 103.62). This indicates that the control compound, 2-Thio-6-Azauridine, does not consistently reach 50 or 95% antiviral reduction levels.

The mean Antiviral Inhibitory Concentration 25% (IC₂₅) was 21.40 μ g/ml (SD \pm 36.0). The median IC₂₅ value was 8.40 μ g/ml (range 0 - 19.7 μ g/ml). The mean Antiviral Inhibitory Concentration 50% (IC₅₀) was 5.60 μ g/ml (SD \pm 15.60). The median IC₅₀ value was 0 μ g/ml (range = 0 - 68.4 μ g/ml). The mean Antiviral Inhibitory Concentration 95% (IC₉₅) could not be attained with 2-Thio-6-Azauridine versus Venezuelan Equine Encephalomyelitis Virus. This discrepancy indicates that the control compound 2-Thio-6-Azauridine does not consistently reach 50 or 95% reduction levels.

The average maximum antiviral inhibitory level of 42 2-Thio-6-Azauridine tests (Figure 25-B) was reached at 100 μ g/ml of the compound with 30% antiviral effect. Further increase of the drug concentration does not improve its antiviral activity. Maximum antiviral effect (~36%) was found with a simultaneous ~50% cytotoxic suppression. Above the 100 μ g/ml concentration the antiviral protection levels off to ~20% reduction level at 320 μ g/ml while simultaneously the 2-Thio-6-Azauridine becomes maximally toxic (~60%).

2-THIO-6-AZAURIDINE - VE VIRUS

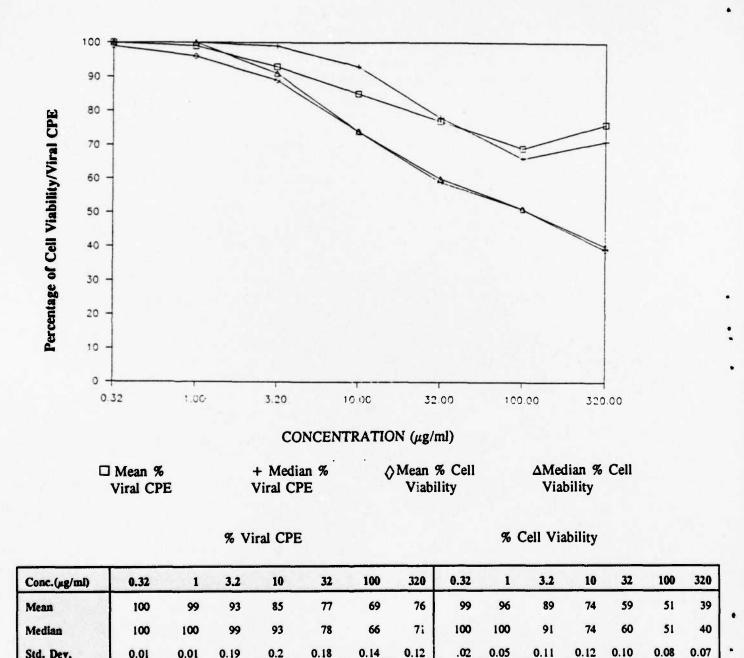


Figure 25-B Average Antiviral and Cytotoxicity Values for 42 Positive Control Compound Tests

0.14

0.12

.02

0.05

0.11

0.12

0.10

0.07

0.18

Std. Dev.

0.01

0.01

0.19

0.2

4.1.5.1.2 <u>Maximum Antiviral Effect of 2-Thio-6-Azauridine vs VE Virus (VE)</u>: Since the metabolic activity of the cells was an unknown function during the testing period, it was monitored indirectly by measuring the maximum antiviral effect of the control compound 2-Thio-6-Azauridine. This demonstrated the amount of infectious virus produced by the cells (Maximum Percent CPE).

A bar graph scatter plot (Figure 26-B) depicts the distribution of the maximum antiviral reduction values of all 42 control compound assays for 2-Thio-6-Azauridine. The results indicate that the average maximum antiviral reduction obtained with the present SOP is around 36% (SD \pm 20.00) reduction levels. The maximum reduction levels vary from 9 - 98% but remain quite consistently around the median of 35%. The assay control values give a relatively broad, bell-shaped distribution curve. This indicates quite a consistent day-to-day performance of the control compound in the VE-MTT assay.

Recommendations:

Based upon the data obtained in parallel studies with Selenazofurin, we do not recommend that 2-Thio-6-Azauridine (AVS-6724) be used as a second control compound against the VE virus. Its overall performance is not as good as that obtained with the present control compound, Selenazofurin. 2-Thio-6-Azauridine is cytotoxic at lower drug concentrations than the observed antiviral effect (see Figure 25-B).

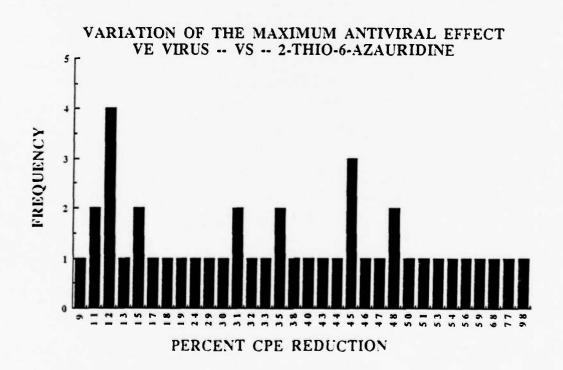


Figure 26-B

Maximum Antiviral CPE Reduction (%).

Summary of 42 Control Tests.

4.1.5.1.3 Cellular Cytotoxicity of 2-Thio-6-Azauridine vs VE Virus:

VE-Control Compound-Cytotoxicity Performance: The 42 cytotoxicity values of the positive control compound 2-Thio-6-Azauridine are also very consistent. The mean cell Toxic Concentration 25% (TC₂₅) was 13.10 μ g/ml (SD \pm 9.50) and the median was 9.80 μ g/ml (range of <1.58 - 38.50 μ g/ml). The mean cell Toxic Concentration 50% (TC₅₀) value was 89.30 μ g/ml (SD \pm 45.20) and the median was 100 μ g/ml (range of 9.65 - 247.00 μ g/ml). The mean cell Toxic Concentration 95% (TC₉₅) value cannot be attained with 2-Thio-6-Azauridine versus Venezuelan Equine Encephalomyelitis Virus.

As can be seen from Figure 25-B, the toxicity starts to become measurable above the concentration of 1.0 μ g/ml and the maximum toxicity has not been reached at 320 μ g/ml.

When the cytotoxicity reaches around 50% (100 μ g/ml), the control compound (2-Thio-6-Azauridine) has not reached its maximum toxicity. After 100 μ g/ml the antiviral protection of 2-Thio-6-Azauridine starts to decrease down to ~20%. 2-Thio-6-Azauridine becomes maximally toxic at 320 μ g/ml. The highest 2-Thio-6-Azauridine concentration tested in these assays was 320 μ g/ml.

2-Thio-6-Azauridine has a definite cytotoxic suppression on cellular metabolism and growth. The TC_{25} and TC_{50} toxicity can be achieved with relative consistency at 320 μ g/ml. As can be seen from Figure 25-B, 2-Thio-6-Azauridine, is cytotoxic at lower drug concentrations than its antiviral activity.

4.1.5.1.4 <u>VE-Assay Plate Ouality Controls: Cell Load and Virus Load Parameters (2-Thio-6-Azauridine):</u> The MTT assay is fundamentally dependent upon the quality of the assay plates. Our large-scale antiviral testing is dependent upon the uniformity of the test plates produced for the daily assays. Equal loads of cell load and virus load as well as the consistent performance of the reagents used daily was monitored. A sample of the plate variation control for the period of November, 1989 through January, 1991 is presented in Figures 27-B, 28-B, and 29-B.

<u>VE-Control Compound-Cell Load Performance</u>: A bar graph scatter plot of the mean cell control (O.D. reading) of 42 control assays is plotted in Figure 27-B. The results indicate that the cell O.D. readings reached a mean 1.140 (SD \pm 0.130) with a median of 1.150 (range of 0.800 - 1.430). This indicates that a uniform and equal number (18,000 cells/well) of cells are being loaded into every well in the 96-well plate during the day-to-day operation. The cells reduced MTT to formazan giving maximum blue color uniformly and consistently.

<u>VE-Control Compound-Virus Load Performance</u>: A bar graph scatter plot of the mean virus load O.D. readings of the 42 control assays is presented in Figure 28-B. The results indicate that the average virus load O.D. reading is 0.050 (SD \pm 0.090) with a median of 0.020 (range of 0.030 - 0.470). This demonstrates that a good cell destruction is taking place and a uniform load of virus (32 TCID₅₀) is administered on the cell monolayer with very consistent viral CPE results.

<u>VE-Control Compound-Assay Differential Performance:</u> A bar graph scatter plot of the mean O.D. differential values of the 42 control assays is provided in Figure 29-B. The results indicate that the average differential O.D. reading is 1.100 (SD \pm 0.150) with a median of 1.100 (range 0.751 - 1.415). The single bell-shaped curve is reasonably sharp and uniform. This reflects that the assays are executed consistently and are repeatable during day-to-day operation with close to 90% measurement accuracy.

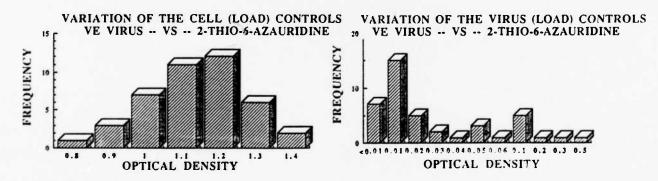


Figure 27-B

Figure 28-B

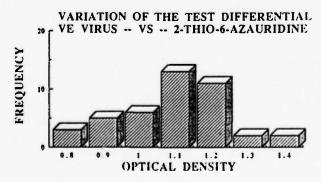


Figure 29-B

4.1.5.2 VE-Antiviral Activity Results:

New Drugs with 95% Antiviral Reduction Levels: Out of the 6067 actual single drug tests, 15 new compounds demonstrated excellent antiviral activity, having antiviral reduction values of equal to or better than 95%. This represents around 0.2% of the test compounds being active at this excellent reduction level. These compounds are summarized in Table 13 according to the highest Total Antiviral Index (TAI). Compound AVS-4051 demonstrated the best in vitro promise, having a TAI of 87% and Selectivity Index (SI) of 313. Six other compounds, AVS-2318, 5906, 2506, 1841, 2960 and 5072, demonstrated good antiviral activity with TAI's of 50, 35,33,32,28 and 2790, respectively, and SI values of 19, 8.9, 9.4, 0.14, 0.31, and 5.0.

It is worthwhile to note that compounds received in shipment number 62 were mostly colored (Table 13). Therefore those compounds appearing in the 95% active category from shipment number 62 should be interpreted with caution, since colored compounds create false positive readings with the MTT assay.

Table 13 $\hbox{AVS Compounds Active Against Venezuelan Equine Encephalomyelitis Virus (VE) at λI_{95} Level }$

Virus	AVS No.	Ship- ment#		Diff- rntl.		IC 95		TC 95		AI 95	SI	TAI
VE	4051	37	09/11/89	1.008	<	0.03	>	10.00	>	312.50 >	312.50 >	86.80
VE	2318	13	09/11/89	0.848		0.94	>	10.00	>	10.68 >	19.40 >	49.94
VE	5906	61	11/03/89			9.42		96.60		10.25	8.87	34.69
VE	2506	64	03/13/90	1.049		2.83	>	320.00	>	113.17 >	9.35 >	33.39
VE	1841	33	10/06/89	0.598		288.00	>	320.00	>	1.11	0.14 >	31.88
VE	2960	27	10/05/90	1.095		875.00	>	1000.00	>	1.14	0.31 >	28.46
VE	5072	48	03/03/89	0.956		264.00	>	320.00	>	1.21 >	4.77 >	27.35
VE	6071	62	01/09/90	1.053		9.59		88.20		9.19	4.04	19.59
VE	7085	72	10/26/90			96.80	>	100.00	>	1.03 >	3.93 >	18.41
VE	6532	66	05/25/90	0.912		94.40	>	320.00	>	3.39	0.01	17.29
VE	8513	76	12/21/90	1.069		941.00		3090.00		3.28	2.84 >	14.49
VE	5121	56	11/02/90	1.165		300.00	>	320.00	>	1.07 >	1.88 >	13.87
VE	2453	18	09/08/89	0.889		9.85	>	32.00	>	3.25	2.64	12.09
VE	2581	65	04/13/90			309.00	>	320.00	>	1.04 >	1.79 >	7.94
VE	3171	29	05/15/89			97.00	>	100.00	>	1.03	0.26 >	6.74

New Drugs with 50% Antiviral Reduction Levels: Out of the 6067 actual single drug tests, 156 new compounds demonstrated good antiviral activity, having antiviral reduction values equal to or better than 50%. This represents around 2.6% of the test compounds being active at this good antiviral reduction level. These compounds are summarized in Table 14 according to the highest Total Antiviral Index (TAI). AVS-5580 and 0111 demonstrated the best TAI's of 47% and SI's of 1780 and 15, respectively. Seven other compounds demonstrated moderate antiviral activity, having TAI's that ranged from 31 - 40% and SI's from 3 - 29. The rest (147 compounds) showed marginal antiviral activity with TAI's that ranged from <1 to 13% and SI's from <1 to 2.

It is worthwhile to note (Table 14) that compounds received in shipment number 62 were mostly colored. Therefore those compounds appearing in the 50% active category from shipment number 62 should be interpreted with caution, since colored compounds create false positive readings with the MTT assay.

Table 14

AVS Compounds Active Against Venezuelan Equine Encephalomyelitis Virus (VE) at AI₅₀ Level

	AVS	Ship-	Test	Diff-													
Virus		ment#		rnt1.		IC	50		TC 50		AI	50		SI		TAI	
VE	5580	54	05/03/89	1.459		56.	20	>	100000	>	1780.	00	>	1780.0	>	46.70	
	0111	9	12/08/89						1000.00		69.		-			46.69	
	5543	56	08/11/89				20		32.00							39.55	
	6218	62	12/07/90			20.							>			39.49	
VE	0053	64	03/09/90			10.	50	>	320.00							39.34	
	5609	57	06/26/89	0.975			88		320.00							36.18	
VE	5643	57	08/18/89	1.000		37.	70	>	320.00	>		48				35.52	
VE	0217	61	12/01/89	1.169		19.			320.00	>	16.	04		14.66	>	32.97	
	0148	2	08/25/89			0.		>	3.20	>	158.	70		4.80	>	31.26	
	2320	13	09/11/89				22		157.00			80			>	28.83	
	2591	65	04/13/90			84.			924.00			00		6.50		25.69	
	3377	65	04/20/90			29.	30	>				94	>		>		
	2433	17	09/11/89			52.			222.00			25		3.08		23.90	
	7048	69	08/03/90			48.	40		210.00		4.	34				23.46	
	6788 1829	67 32	07/13/90 10/06/89	1.189		27.	90	>	1000.00	>	11.	68				22.89	
	2503	21	09/08/89				68		10.00		14.		•			21.97	
	4742	44	09/05/89			19.						58				21.61	
	5497	53	04/17/89			354.						90				20.90	
	0124	64	03/09/90			25.						81				20.48	
	5040	45	01/05/90			72.			899.00			34		4.17		20.36	
	5714	58	10/09/89			1.			6.38			28		3.04		20.06	
	4871	46	01/16/89			50.		>				29			>	19.83	
	5197	58	10/09/89			21.			164.00		7.	46		4.04		19.73	
VE	6489	66	05/18/90			2.	70		66.90		24.	77		16.26	>	19.55	
VE	5724	58	10/09/89			204.	00	>	1000.00	>	4.	90	>	4.90	>	19.40	
VE	2573	65	04/13/90			220.	00	>	1000.00	>	4.	55				19.00	
	7116	70	09/14/90					>	1000.00			90	>				
	5384	54	05/01/89			17.			66.70			73				18.03	
	2590	19	09/08/89			211.			320.00			52	>		>	17.85	
	5119	56	08/07/89			24.		>	100.00			80		3.08		17.35	
	5138	57	08/18/89			179.			711.00			97				17.32	
	3802	67	07/13/90				23	>	10.00			52				17.25	
	5198 2275	58 12	10/09/89			19. 51.			84.90 284.00			41 55				17.11 16.91	
	5383	54	09/11/89 05/01/89			64.			211.00			27				16.70	
	6906	69	07/19/90			17.			60.90			54				16.44	
	6029	61	11/10/89			86.			320.00			70	•			16.40	
	0230	1	08/25/89			20.			100.00			98				16.03	
	5485	53	12/15/89			7.			32.00			05				15.90	
	1159	52	08/25/89			64.			283.00			41		2.99		15.49	
	6837	68	06/15/90			85.	00		215.00			53		1.85		15.32	
VE	8318	76	12/14/90	0.465		592.	00		2240.00			78				15.27	
	2631	65	07/06/90	1.033		193.			732.00			79				15.16	
	5997	61	11/10/89		<	1.	00		57.50	>	57.	50	>		>	14.84	
VE	230	1	12/12/88	1.299			67	>	100.00		11.			2.54		13.79	
	5175	58	10/09/89			20.			59.90			94			>	13.52	
	3378	65	04/20/90			17.					17.			0.44		13.41	
	7087	72	10/26/90			20.		>				83				13.41	
	8397	76	01/08/91			667.			1690.00			54			>	13.41	
	4770	44	11/08/88			59.			210.00			50		2.59		13.39	
	8601 4769	76 44	01/11/91			24.			124.00			05		2.86		12.94	
	3611	32	11/08/88			59.	91		205.00		13.	45		2.49		12.90	
	6315	63	10/05/90				38		25.70 21.10			31				12.83	
	6482	66	05/18/90			59.			198.00			32		2.29		12.72	
	5385	54	05/01/89			26.			65.60			50			>	12.33	
	6521	66	05/25/90				09	>	320.00		39.			27.38			
	7031	69	08/03/90			76.			320.00			16		2.86		12.26	
	6943	69	09/07/90			80.			320.00			97		2.61		12.16	
	3584	32	09/11/89			72.			285.00			93		2.45		11.91	
	Washington .						1 10 1				-						

Table 14 (Cont'd)

	0.000	St. 1			- 111				
Virus	AVS	Ship- ment#		Diff- rntl.	IC 50	TC 50	AT 50	er ma	T
ATLUM	NO.	mence	DACE	rnei.	10 50	10 50	AI 50	SI TA	Ļ
VE	5691	57	08/18/89	1.091	200.00	734.00	3.67	2.35 11.53	2
	1019	28	08/25/89		0.55 >	1.00 >	1.83 >	1.83 > 11.4	
	3392	65	07/20/90		179.00 >		1.79 >	1.79 > 11.39	
	0646	2	08/25/89	0.866	0.79 >		4.06	1.18 > 11.20	
VE	2585	65	04/13/90	1.063	189.00 >		1.70 >	1.70 > 11.1	
	6311	63	02/16/90	1.298	72.80 >	320.00 >	4.40	1.06 > 11.0	
	4892	46	01/17/89		273.00 >	320.00 >	1.17	0.24 > 10.89	
	1850	32	10/06/89		77.00 >		1.30 >	1.30 > 10.8	
	5457	54	05/02/89		15.70	48.40	3.08	0.48 > 10.84	
	2582	65	04/13/90			1000.00 >	1.67 >	1.67 > 10.70	
	7086 7047	72 69	10/26/90		283.00 >	100.00 >	4.61	0.64 > 10.69	
	5176	58	08/03/90 10/09/89		79.20	320.00 > 200.00	1.13 2.52	0.89 > 10.60 1.69 10.40	
	7017	69	07/27/90		197.00 >	320.00 >	1.62 >	1.62 > 10.3	
	8511	76	12/21/90			3200.00 >	1.35 >	1.35 > 10.34	
	4919	46	01/31/89		97.00	310.00	3.20	1.67 10.3	
	4747	44	12/15/89		32.00	73.50	2.30	1.65 > 10.10	
VE	0206	4	12/08/89		268.00	841.00	3.14	1.79 9.99	
VE	5998	61	02/09/90		4.97	23.30	4.69	1.67 9.94	4
11-	7083	72	10/26/90	1.152	93.50 >			2.06 > 9.89	9
	8212	75	11/30/90			3200.00 >		1.34 > 9.72	
	2716	22	09/08/89		8.43 >	10.00 >	1.19 >	1.19 > 9.60	
	4768	44	11/08/88		64.90	194.00	2.98	2.01 9.5	
	2600	65	04/13/90			1000.00 >	2.02	1.68 > 9.3	
	8315 4852	76 48	12/14/90		729.00	1960.00	2.70	1.85 > 9.33 0.08 9.33	
	8396	76	02/21/89 01/08/91		47.80 1000.00	9.55 1400.00	0.20 1.40	0.08 9.32 0.60 > 9.28	_
	7009	69	07/27/90	1.107		320.00 >		1.69 > 9.19	
	7032	69	08/03/90		273.00 >		1.17 >	1.17 > 9.14	
1	5058	48	01/05/90			1000.00 >	1.48	1.48 > 9.13	
	8398	76	01/08/91		661.00	1990.00	3.01	0.45 > 8.64	
	3935	65	07/06/90		69.40	175.00	2.52	1.48 8.59	
	5531	56	08/11/89	1.293	80.20	291.00	3.63	2.14 8.50	
	6986	68	08/10/90		227.00	520.00	2.29	1.25 8.20	
	1215	27	10/06/89		100.00	268.00	2.68	1.37 8.03	
	6321	63	02/16/90		18.40	50.90	2.77	1.30 7.93	
	5905	61	11/03/89		7.41	19.20	2.60	1.68 7.83	
	3038 2989	28 27	09/08/89 04/14/89		0.70 > 27.10	3.20 > 63.40	4.56 2.34	0.41 > 7.76 1.06 7.69	
	8400	76	01/08/91			3200.00 >	1.00	0.40 > 7.59	
	7016	69	09/07/90		216.00	576.00	2.67	1.65 > 7.34	
	7403	70	09/28/90			3200.00 >	1.08 >	1.08 > 7.34	
	0346	2	05/09/89	0.879	2.76	17.70	6.42	0.83 > 7.2	
	6994	68	06/22/90		203.00 >	320.00 >	1.58 >	1.58 > 7.13	
VE	3688	32	11/10/88		287.00 >	320.00 >	1.11 >	1.11 > 6.84	
	2572	65	04/13/90		221.00 >	320.00 >	1.45 >	1.45 > 6.83	
	5541	56	06/16/89	1.276	100.00	26.50	0.27	0.07 > 6.73	
	7444	73	10/26/90			3200.00 >	1.49	1.49 > 6.53	
	5379	54	05/01/89		6.84	20.00	2.92	0.41 > 6.48	
	7386	70	09/28/90			3200.00 >	1.52 > 5.12 <	1.52 > 6.34	
	5854 2902	60 26	10/20/89 04/14/89		62.60 > 70.00	320.00 > 155.00	2.21	0.02 > 6.22 0.89 6.14	
	8270	76	12/14/90			3200.00 >	1.30	0.89 6.14 0.84 > 5.90	
	6140	62	01/04/90		7.56	22.60	2.99	1.17 5.7	
	5094	56	06/19/89		73.10	156.00	2.13	1.14 4.5	
	5495	53	04/17/89		24.90	8.76	0.35	0.20 > 4.46	
	2543	21	09/08/89		30.00	73.00	2.44	1.46 4.4	
	2544	20	12/20/88		2.60	6.20	2.38	1.28 4.34	
	6309	63	02/16/90	1.170	198.00 >	320.00 >	1.62	1.12 > 4.2	7
	8696	76	01/25/91		271.00 >	320.00 >	1.18 >	1.18 > 4.23	
	5489	66	05/11/90		305.00 >	320.00 >	1.05 >	1.05 > 3.93	
The second second	5142	57	07/10/89		81.10	161.00	1.98	1.04 3.3	
	3559	31	03/28/89	The state of the s	71.40	193.00	2.71	1.20 3.2	
VE	8374	76	12/14/90	1.1/4	957.00	1780.00	1.86	1.02 3.14	4

Table 14 (Cont'd)

***	AVS	Ship-		Diff-		_	70.50					m> -
Virus	NO.	ment#	Date	rntl.	IC 5	U	TC 50		AI 50	SI		TAI
VE	8241	75	11/20/90	1.074	1990.00	0 >	3200.00	>	1.61	0.35	>	3.12
VE	6412	66	05/11/90		91.60	0	157.00		1.71	0.90		3.06
VE	5907	61	02/02/90		458.00	0	100.00		0.22	0.10	>	2.91
VE	6477	66	05/18/90		24.90	0	57.90		2.33	0.56		2.91
VE	6659	64	04/06/90		261.00	0 >	320.00	>	1.23	0.82	>	2.83
VE	8395	76	01/08/91		2780.00	0 >	3200.00	>	1.15	0.29	>	2.83
VE	8221	75	11/20/90	1.108	19.90	0	62.20		3.13	< 0.50		2.79
VE	7059	72	08/28/90	1.294	179.00	0	64.00		0.36	1.79	>	2.53
VE	7460	73	11/02/90		26.50	0 >	32.00	>	1.21	0.79	>	2.29
VE	4409	44	12/13/88		252.00	0 >	320.00	>	1.27	0.87	>	2.15
VE	3160	29	04/28/89		23.00	0 >	32.00	>	1.39	0.27	>	1.91
VE	0113	1	05/22/89	0.623	92.50	o >	320.00	>	3.46	0.33	>	1.87
VE	7911	75	11/09/90		2390.00	0 >	3200.00	>	1.34	0.27	>	1.87
VE	6993	68	06/22/90		296.00	>	320.00	>	1.08	0.74	>	1.78
VE	5070	48	03/03/89	0.958	232.00	0	97.00		0.42	0.26	>	1.63
VE	6791	67	07/13/90		24.50	0	4.35		0.18	0.10		1.54
VE	6070	62	01/09/90		8.56	5	13.70		1.60	0.47		1.16
VE	5393	54	05/02/89	1.045	9.4	5	2.91		0.31	0.20	>	1.09
VE	3560	31	03/28/89	1.022	287.00	0 >	320.00	>	1.12	0.40	>	0.75
VE	3164	29	04/28/89	0.937	32.00	0	37.10		1.16	0.30	>	0.68
VE	6739	64	03/09/90	1.158	249.00	0 >	320.00	>	1.28	0.27	>	0.63
VE	3322	65	04/20/90		10.00	0	9.69		0.97	0.58	>	0.60
VE	6310	63	02/16/90	1.298	32.00	0	31.00		0.97	0.09		0.58
VE	3980	36	09/11/89	1.141	100.00	0 >	100.00	>	1.00	0.32	>	0.40
VE	4998	51	03/14/89	0.983	249.00	0 >	320.00	>	1.28	0.11		0.36
VE	4999	51	03/14/89		88.50	0	2.48		0.03	0.02	>	0.22
VE	5690	57	07/17/89	0.883	306.00	0 >	320.00	>	1.05	0.21		0.10
VE	6373	63	03/02/90	1.122	278.00	0	308.00		1.11	0.24	>	0.08
VE	2430	17	03/17/89		24.80	0	59.30		2.39	1.21	>	0.00
VE	5528	56	06/12/89		7.84	4 >	320.00	>	40.80	0.86	> 1	5.38
VE	5531	56	06/12/89	1.225	240.00	0 >	320.00	>	1.33	0.97	>	2.55

New Drugs with 25% Antiviral Reduction Levels: Of the 6067 actual single drug tests, 356 new compounds demonstrated moderate antiviral activity, having antiviral reduction values equal to or better than 25%. This represents around 6% of the test compounds being active at this marginal antiviral reduction level.

In general, when compared to the 95% and 50% antiviral activity categories, the compounds in this (25%) category do not appear to have any significant antiviral promise and probably do not need to be presently confirmed any further.

4.1.5.3 Confirmatory Assays:

Some of the compounds were sent to us in more than one separate drug shipment. These compounds were tested more than once. Data from the confirmatory assays are summarized in Table 15. If a compound showed $\geq 50\%$ reduction in CPE during this contract period, then it was considered a candidate for confirmatory testing. The confirmatory tests are from active compounds picked up by both the VR and MTT assay testing. Out of 131 confirmatory tests, 85 compounds were confirmed active during this reporting period and the remaining 46 compounds gave conflicting results. The criteria for activity is that the confirmatory test has to show $\geq 25\%$ reduction in CPE. Failure to confirm the activity in these compounds was probably due to differences during the assay conditions:

- 1) In confirmatory assays the concentration range is adjusted to a more accurate semilog scale to maximize the TAI window.
- 2) Differences in the "differential" of the two runs can cause the compound to read positive or negative, falsely. The variability in the differential can cause false positive or negative bias to be introduced into the calculations, thus reflecting the variability in the maximum activity of the compound.
- 3) The metabolic rate, cell and virus load/well, age, and passage number of the cells may cause the above observed variability in the confirmatory results.
- 4) Problems associated with stability and storage of the compound (i.e., different lot numbers, solubility, light sensitivity, hygroscopic, etc.).
- 5) Problems associated with technical execution of large numbers of plates by different technicians.

During this reporting period the overall confirmatory rate against VE was 65%. The conflicting results should be retested at a later date based on the availability of the compound.

4.1.5.4 Recommendations of VE-Actives Based Upon the *In Vitro* Results with MTT Assay (Vero Cells)

Based upon the in vitro results with the MTT assay (Vero cells) we recommend the following:

- 1) Compounds with the highest TAI in the 95% activity category that have confirmed results with the exception of "colored" compounds should receive the highest priority for further profile studies and *in vivo* animal testing.
- 2) Compounds with the highest TAI in the 50% activity category that have confirmed results with the exception of "colored" compounds should receive the next highest priority for further profile studies and *in vivo* animal testing.

Table 15

Confirmatory Assays for Compounds Active Against Venezuelan Equine Encephalomyelitis (VE)

∢ ∪⊢	* * *	• • • •	+ +	* * *	* * *		• • • •	++ •	+ +++	+ + +	• •
Assay Type	### ###		EE	FFF	FFF	MATTER TE					FF
TAI	39.34 20.28 9.08	6.28 18.41 29.87 46.69	1.87	20.48 9.64 11.24	31.26 0.00 0.07	0.00 9.03 1.98 1.94	10.09 32.97 11.20 0.00	9.20	8.01 21.97 2.99	3.73	10.87
13	28.68 × 2.92 × 0.00 ×	0.84 2.22 × 4.58 × 15.22 ×	0.33 ×	3.28 × 1.26 × 2.15 ×	4.80 v	2.40 0.00 1.73 0.00 v	1.18	0.94	0.00 \$ 11.51 \$ 0.00 \$	0.00	1.30 >
A1 95	0.00	0.000	0.0	888	8000	0.0000	0.00	0.00	0.00	0.00	0.00
75 %	320.00 320.00 320.00	320.00 100.00 1000.00	320.00	320.00 1000.00 1000.00	3.20 3.20 10.00	320.00 320.00 1000.00 1000.00	320.00 320.00 3.20 3.20	100.00	320.00 320.00 320.00	320.00 > 966.00 943.00	1000.00
10.95	0.00	0.000	0.00	90.00	0.00	0.0000	0.00	0.00		288.00 v 0.00 0.00	0.00
AI 50	30.42 21.27 0.00	2.52 4.76 50.57 69.19	3.46	12.81 4.19 5.87	158.70 0.00 0.00	2.60 3.00 3.14 0.00	0.00 16.04 0.00	1.83	0.00	6.36 0.00 0.00	1.30
10 50	320.00 > 146.00 24.90	74.50 100.00 > 1000.00 >	320.00 > 100.00	320.00 > 93.80 141.00	3.20 > 0.94	320.00 × 320.00 × 611.00 841.00 272.00	320.00 320.00 > 3.20 > 0.80	1.00 > 47.70	320.00 320.00 1000.00	320.00 × 660.00 433.00	100.00 > 348.00
10 50	10.50 × 6.84 0.00	29.50 21.00 × 19.80 × 14.50 ×	92.50 > 0.00 >	25.00 × 22.40 × 24.00	0.00	6.00 v 0.00 v 0.00 v 0.00 0.00 0.00 0.00	0.00 19.90 v	2.33	0.00 v 27.80 v 0.00 v 0.00 v	19.60 v 0.00 0.00	7.00 > 0.00
AI 25	60.82 11.83 3.67	2.30 5.08 7.57 27.43	1.05	3.29	7.36	3.20 0.99 1.57 3.77	3.53 23.09 2.13 0.62	1.83	0.00	1.86	2.35
10 25	302.00 20.00 9.06	24.90 46.60 90.60 220.00	30.80	82.00 28.30 51.70	0.10	320.00 - 198.00 272.00 479.00 124.00	57.10 293.00 0.93 0.31	× 1.00 × 2.20 × 83.60	> 320.00 > 320.00 > 320.00 > 927.00	2.79 490.00 238.00	> 100.00 > 131.00
10 25	4.96 1.69 2.47	10.80 9.16 12.00 8.02	48.40	15.50 12.40 14.40	0.01	- 100.00 199.00 173.00 127.00 80.30	16.10 12.70 0.44 0.50	1.20	25.30	4.39 264.00 0.00	42.50
Diff.	1.183 1.521 1.064	0.956 0.761 1.138 1.145	0.623	1.158 1.391 0.966	0.795	NA 0.903 1.114 1.272 0.958	1.004 1.169 0.866 0.964	0.781	1.042	0.598 1.003	0.716
Test Pit Date #	03/09/90 UGB 05/04/90 VTM 10/05/90 29F	05/09/89 Q0G 08/25/89 RAC 12/08/89 SUZ 12/08/89 STT	05/22/89 Q5K 08/25/89 RAC	03/09/90 UGF 05/04/90 VTO 07/20/90 XKH	08/25/89 RAD 06/01/90 WES 10/05/90 29G	05/14/87 05/09/89 00H 12/08/89 SS9 12/08/89 SS0 06/01/90 WE5	04/17/89 PLV 12/01/89 SQU 08/25/89 RAH 06/01/90 WE6		10/06/89 RTW 04/03/89 P8V 10/06/89 RU2 06/01/90 WHO	10/06/89 RU3 01/19/90 TG4 04/13/90 V98	10/06/89 RU4 06/01/90 WHO
Ship- ment		88	58	222	191	_	m- h	28 08 28 03 27 04		33 10 62 01 65 04	32 10 67 06
AVS S	** VIRUS VE 0053 64 0053 64 0053 64	0111 0111 0111 0	0113 1	0124 6 0124 6 0124 6	0148 2 0148 6 0148 2	0206 0206 0206 0206 0206 0206 0206	0217 3; 0217 6; 0646 2 0646 6;	1019 2 1019 2 1215 2		1841 3 1841 6 1841 6	1850 3 1850 6

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Assay				#### #####	TIM	FFFF		ĒĒ	ĒĒ	ĒĒ	ĒĒ	ĒĒ
IAI	2.62 16.91 0.31	0.42 49.94 0.10 14.40	13.05 28.83 15.87 17.17 15.07	3.43	12.09 1.57 0.41	2.33 21.71 0.00 0.67 0.01	13.28 33.39 14.16	4.34	1.24	0.00	17.85	8.35
15	0.00	0.25 v 19.40 v 0.00 v 2.03 v	0.00 6.39 5.41 2.78 2.98	3.08	2.64 0.00 0.00	0.00 0.00 0.00 0.00 0.00	2.81 > 9.35 > 2.90 >	1.30	3.72 >	0.00	1.52 >	2.11 > 6.50
A1 95		0.00 83.00 0.00	00000	0.00.0	3.25	00000	0.00	0.00	0.00	0.00	0.00	0.00
70.95	100.00 320.00 320.00	6.6.6 8.8.8 \$	100.00 320.00 320.00 320.00	100.00 320.00 320.00	32.00 × 89.90 71.70	3.20 10.00 10.00 32.00	10.00 320.00 >	27.00	320.00	1000.00	320.00	320.00
10.95	0.00	0.00.00	00000	00000	9.85 v 0.00 0.00	00000	0.00 × 0.00 × 0.00 ×	0.00	0.00	0.00	0.00	0.00
A1 50		1.44 19.40 0.00 4.87	0.00 21.80 18.55 8.91	1.14	4.13 0.00 0.00	7.82 0.00 0.00 0.00	4.42 80.57 19.78	0.00	0.00	1.67	1.52 2.26	11.00
16 50		7.00.01 70.00 70.00 70.00	22.70 157.00 28.50 54.70 83.00	100.00 × 222.00 × 320.00 × 188.00	20.70 25.30 19.70	5.6.3 5.6.3 5.6.3 5.6.3	10.00 × 80.60 × 19.80 ×	6.20	320.00	1000.00	320.00 >	320.00 >
10 50		1.00	0.00 7.22 1.53 6.14 8.27	87.90 × 52.30 × 222.00 × 0.00	5.02 0.00 0.00	0.00 88.00 0.00 0.00 0.00 0.00	2.26 > 1.00	2.60	0.00 > 220.00 >	0.00 >	211.00 > 443.00 >	152.00 >
A1 25		0.69 312.50 0.56 3.45	13.10 46.17 8.30 17.04 13.41	21.78 21.78 1.77 0.00	3.85 0.80 1.27	1.51 10.38 0.74 1.20 0.26	6.18 9.35 < 2.90 <	2.10	0.00	0.00	43.66	4.53
10.25		0.25 10.00 v 0.31	8.93 46.20 > 8.30 > 17.00 >	161.00 236.00 75.10	13.30 17.60 11.60	0.28 4.17 0.54 2.18 1.30	6.37 9.35 > 2.90 >	3.30	320.00	1000.00	320.00 > 852.00	547.00
10.25		0.36 0.03 v	6.00 1.00 1.00 1.00 1.00	51.40 × 7.40 133.00 0.00	3.45 22.20 9.16	0.18 0.73 1.81 4.97	1.03	1.60	0.00 >	342.00 >	7.33 >	51.90
Diff.	1.050 0.944 0.987	1.040 0.848 0.987 1.079	1.067 0.848 1.171 1.019	0.950 0.631 1.171 1.026	0.889 0.995 1.260	1.059 1.216 1.009 1.139 1.107	1.020	1.115	1.058	0.951	1.069	1.069
£ **		X E E X	P44 KEI XX7	P58 V90 XA8	RGS UJ7	766 KG XGC XGC XGC XGC XGC XGC XGC XGC XGC XG	RG7 UJ8 VTY	07S P92	0.13 V9E	RGB V9G	RGC V9H	RGC
Test	288	03/17/89 09/11/89 06/01/90 07/13/90	03/17/89 09/11/89 04/13/90 07/06/90	03/17/89 09/11/89 04/13/90 07/06/90	09/08/89 03/13/90 05/04/90	03/31/89 09/08/89 06/01/90 07/13/90 10/05/90	09/08/89 03/13/90 05/04/90	12/20/88 03/21/89	06/26/89	09/08/89	09/08/89	09/08/89
Ship-	21 27	11 79	£ £ 8 8 8	17 65 65	522	22 67 67	222	22	57	59	19 65	65
AVS		2318 2318 2318 2318	2320 2320 2320 2320	2433 2433 2433 2433	2453 2453 2453	2503 2503 2503 2503 2503	2506 2506 2506	7752	2573 2573	2882	2590	2591

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Assay Type	THE T	E E	H H		EEEE			FE				H			
TAI	13.37 10.00 15.16	6.03	6.14	2.64 6.82 28.46	 180.00	0.00	6.74 9.87	17.94	24.94	6.35 8.17 11.91	12.83 0.00 0.00	9.00	0.18 12.01 0.74 17.25	3.04	
5	1.77 • 1.66 • 2.63 •	1.27 >	0.00	4.34 v 0.15 v 0.31 v	0.0.0	0.00	0.26 >	2.40 > 0.00	0.00	0.76 > 0.00 > 2.45	2.49 > 0.00	1.11	0.00 1.33 v 0.00 2.58 v	0.00	
AI 95	0.00	0.00	0.00	1.16 > 1.16 > 1.14	8888	0.00	1.03	0.00	0.00	0.00	0.00	0.00 0.00	9000	0.00	
10 95	320.00 320.00 1000.00	93.60	322.00 100.00	320.00 3 20.00 1 000.00 3	3.20 3.20 320.00 10.00	320.00 10.00 3.20	100.00	320.00	320.00 320.00	100.00 320.00 320.00	320.00 320.00 1000.00	320.00 979.00	32.00 10.00 10.00	299.00 303.00 306.00	
10.98	0.00	0.00	0.00	276.00 × 0.00 × 875.00 ×	00000	0.00	97.00 >	0.00	0.00	0.00	0.00	9.00	00000	888	
AI 50	3.5%	3.15	2.21	4.34 1.91 11.87	0.00	0.00	2.08	0.00	10.94	1.24 0.00 3.93	13.40 0.00 0.00	0.00	0.00 52.25 0.00 42.52	0.00	
1C 50	320.00 × 320.00 × 732.00	8.40	155.00	320.00 > 320.00 > 1000.00 >	1.80 2.40 2.60 1.70	1.00	100.001 100.00	320.00 >	320.00 >	100.00 × 320.00 × 285.00	25.70 320.00 1000.00	320.00 >	0.31 10.00 77.0 10.00	87.10 148.00 175.00	
10 50	181.00 > 193.00 > 193.00	2.67	0.00	73.70 > 167.00 > 84.30 >	0.0000	0.00	48.50 >	19.00 >	29.30 \$	80.50 × 0.00 × 72.60	0.00 \$	287.00 > 0.00	0.00	0.00 76.30 69.40	
A1 25	2.85 2.50 4.55	2.43	1.39	8.70 0.34 1.15	0.00 0.00 13.00 13.00	1.00	0.38	0.00	35.19	3.73	0.00	1.91	1.00 2.65 1.16 3.96	0.92	
10 25	320.00 > 508.00 >	3.40	62.20	. 320.00 > 24.70 > 26.30	0.23 1.10 1.50 0.76	1.00	12.40	44.00	320.00 >	61.10 235.00 178.00	4.77 × 320.00 681.00	320.00 > 272.00	. 0.10 0.25 0.21 0.61	41.20 72.30 103.00	
10 25	112.00 > 128.00 > 112.00	1.40	44.90	36.80 × 73.30 × 23.00	2.40 0.30 12.30 5.70	< 1.00 < 0.07 < 0.26	33.00 18.60	0.00	9.09	21.70 62.90 44.10	0.00	168.00 v	0.10 × 0.10 × 0.10 × 0.10 × 0.10	44.60 47.10 39.30	
Diff.	0.716 1.248 1.033	1.214	1.201	1.162	1.039 0.953 0.931 1.371	0.835 0.854	1.131	1.293	1.182	1.169	1.285	0.747	1.165	1.180 1.313 0.977	
= *	8 8 X	RHZ	E I	P12 R14 291	000 021 PE2 R15	PEF PJE R16	SUS OUR	029 R17	VE0	002 PLR RJG	893	OAA SYN	RJS WH7	RJU VKG XAB	
Test	09/08/89 04/20/90 07/06/90	01/23/89	04/14/89	04/14/89 09/08/89 10/05/90	02/07/89 03/10/89 04/07/89 09/08/89	04/10/89 04/14/89 09/08/89	05/15/89 12/08/89	12/07/88 09/08/89	04/20/90	11/03/88 04/17/89 09/11/89	11/03/88 03/09/90 06/01/90	11/10/88	08/04/89 09/11/89 06/01/90 07/13/90	09/11/89 04/27/90 07/06/90	
Ship- ment	95 65 65	92	28	27	33 22	882	22	28	\$ \$	222	32 25	22	328	888	
No.	2631 2631 2631	2716 2716	2902	2960	2980 2980 2980 2980	3038 3038 3038	3171	3315	3377	3584 3584 3584	3611 3611 3611	3688	3802 3802 3802 3802	3935 3935 3935	

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Assay Type			ĒĒ	EE	MTT	ĒĒ	ĒĒ	EE	HH		FFFF			FF
IĀT	86.80 0.00 0.10 0.03	9.62 21.61 0.00	9.53	12.90	13.39	9.32	19.83	10.89	10.31	20.38 0.38	0.00 6.06 9.13 4.35	0.03	27.35 24.06 3.10	4.52
ïs	312.50 v 0.00 0.00 v 0.	0.17 > 0.38 > 0.00	2.01	0.00	2.59	0.00	3.99 >	0.24 >	1.67	0.00 4.17 0.00 >	1.23 \$ 1.48 \$ 0.00	0.00	5.88 v 0.00 v	1.14
AI 95	312.50 0.00 0.00 0.00	0.00	0.00	0.0	0.0	0.00	9.0	0.00	9.0	0.00	9888	0.00	1.21	0.00
76 27	28.73 28.73 1.88	320.00 320.00 320.00	307.00	309.00	309.00	320.00	320.00	320.00	320.00	1000.00 1000.00 3200.00	320.00 1000.00 1000.00 1000.00	320.00 311.00 301.00	320.00 × 1000.00 320.00	304.00
10 95	0.0000	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00.0	0.00	264.00 \$	0.00
AI 50	312.50 < 0.00 0.00 0.00 0.00 0.00	5.52 16.58 0.00	2.98	3.45	3.50	0.20	6.29	0.00	3.20	0.00 12.34 0.00	1.32	0.00	4.77 7.26 0.00	2.13
10 50	10.00 v 6.63 v 7.66 0.63 v 6.46	320.00 > 320.00 > 25.10	194.00	205.00	210.00 165.00	9.55	320.00 >	320.00 > 171.00	310.00 250.00	861.00 899.00 482.00	320.00 1000.00 > 1000.00 > 827.00	97.00 179.00 94.90	320.00 > 1000.00 > 320.00	156.00 98.50
05 21	0.000.00 0.000.00 0.000.00	58.00 > 19.30 > 0.00	%.0° .0°	59.60	9.00	47.80	50.90 > 63.40 >	273.00 > 0.00	97.00	0.00	0.00 × 675.00 × 0.00	232.00 0.00 0.00	67.00 > 138.00 > 0.00 >	0.00
AI 25	312.50 < 0.00 0.00 0.00 0.00 0.00	1.24	2.88	3.44	3.56	0.17	12.50	4.12	3.60	2.23 7.06 1.11	0.83 2.66 3.98 2.30	0.00	7.87 14.95 1.49	1.73
10,25	10.00 × 5.92 5.19 0.41 4.65	9.68 7.32 16.10	130.00	148.00	155.00	3.88	203.00	64.50	162.00	425.00 304.00 137.00	223.00 935.00 1000.0 518.00	59.20 106.00 52.40	320.00 × 810.00 267.00	83.50 59.80
10 25	, 0.00 0.00 0.00 0.00	9.58 5.91 0.00	45.20	43.10	43.50	22.80	16.20	15.70	44.90	191.00 43.00 123.00	269.00 351.00 252.00 225.00	0.00	40.70 > 54.10 179.00	48.40
biff.	1.008 0.964 1.170 1.234 1.102	0.994 0.805 1.234	1.224	1.117	1.117	1.006	0.802	0.894	1.004	1.343 1.095 0.933	1.000 1.350 1.095 0.970	0.958 1.011 1.368	0.956 1.011 0.390	1.410
*	27 29 29 29 29 29 29 29 29 29 29 29 29 29	P8T RKZ Z9L	280	95	8 5	800	OBF SR5	803 101	015 SR6	SS2 111 1WU	SS3 111 1W	PUZ WE	OVJ RUZ RTN	afe R1L
Test	09/11/89 06/01/90 10/05/90 10/05/90	04/03/89 09/15/89 10/05/90	11/08/88 02/16/90	11/08/88 02/16/90	11/08/88 03/09/90	02/21/89 03/09/90	01/16/89 12/01/89	01/17/89	01/31/89 12/01/89	12/08/89 01/05/90 02/09/90	05/23/89 12/08/89 01/05/90 02/09/90	03/03/89 10/02/89 04/27/90	03/03/89 10/02/89 10/06/89	06/19/89 08/07/89
Ship-	2562	333	33	33	22	87	919	99	32	62 45 GABSN	2323	87 87 87	3 9 9	25 25
AVS .	1507 1507 1507 1507 1507 1507	4742	8927	6927	6773	4852 4	1287	4892 4	6167	2040 2040 2040	5058 5058 5058 5058	5070 5070 5070	5072 5072 5072	5094

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Assay	EEE	HHH		FE				T T T		TIM		T I I	ĒĒ	FE
¥	7.86 17.35 4.18	0.30	3.90 17.32 11.49	3.31	13.52	10.00	19.73	0.93	6.48	16.70	18.03 0.00 2.32	12.33	0.00	10.84
3	3.08	0.00	0.00 v 2.88 v 0.00 v	20.0	0.00	0.00	6.04 8.00 8.00	3.01 ×	0.61	2.41 > 0.00	2.72 v 0.00 0.81	1.82 >	0.20	0.48 >
AI 95	0.00	0.00	888	88	888	888	888	0.0	88	88	0.00	88	88	0.00
25 25	320.00 100.00 320.00	300.00 320.00 320.00 >	320.00 1000.00 1000.00	304.00	92.60 96.70 93.60	320.00 317.00 622.00	301.00 307.00 298.00	302.00	3.20	311.00	99.30 97.30 95.10	97.90	320.00 10.00	155.00
10 %	0.00	300.00	0.00	00.0	888	988	888	0.00	90.0	9.0	0.00	0.0	0.00	0.00
AI 50	0.00	0.00	3.97	1.98	0.20	0.00	1.38	1.58	2.92	3.27	3.73 0.00 1.88	2.50	0.31	3.08
1C 50	215.00 100.00 v 71.50	120.00 247.00 320.00 >	320.00 711.00 1000.00	161.00	8.48 59.90 35.80	92.20 200.00 187.00	127.00 164.00 100.00	140.00	3.07	211.00	66.70 57.10 50.70	65.60	10.00	48.40
10 50	0.00 24.50 v 0.00	0.00	178.00	81.10 0.00	0.00	9.50	21.90	88.50 19.30	6.8 0.00	64.60 0.00	17.90 0.00 27.00	26.20	9.45	15.70
AI 25	2.59 7.07 0.00	0.00	1.27	1.67	0.31 4.54 0.56	3.52	0.82 5.99 1.15	0.60	0.0	4.41	5.56 0.00 1.37	3.70	1.87	4.21
70 25	92.70 75.60 50.50	69.80 173.00 320.00 >	. 320.00 > 515.00 · 1000.0 >	84.30	5.54 39.40 8.87	17.30 134.00 106.00	44.40 88.70 66.00	32.00	2.83	156.00 66.00	48.60 34.70 21.80	47.70	1.87 >	5.79
10 25	35.90 10.70 0.00	0.00	251.00 > 110.00 > 273.00 >	50.50 57.20	17.90 8.67 15.90	259.00 38.10 67.70	54.20 14.80 57.30	53.20	3.78	35.30	8.75 0.00 15.90	12.90	4.00 0.00	1.79
Diff.	0.677 1.123 1.397	0.590 1.069 1.165	1.145	1.103	0.703 1.000 1.383	0.703	1.010	1.439	0.943	0.794	0.795	0.795	1.045	1.072
£*	19 9H4 19 81N	19 9HS 19 R10 10 ZHL	9 GPY 19 R7C	90 800 870 870	9 QVE	S GVE	19 QY2 19 RYU 10 Z9H	19 GY2	9 PVH 970 Q	N60 68	19 PVP 19 090 10 VMF	9 690 690	9 PX4	9 PXB
Test Date	06/23/89 08/07/89 04/27/90	06/23/89 08/07/89 11/02/90	07/10/89 08/18/89 04/27/90	07/10/89 08/18/89	07/24/89 10/09/89 04/27/90	07/24/89 10/09/89 04/27/90	07/28/89 10/09/89 10/05/90	07/28/89 10/09/89	05/01/89 06/09/89	05/01/89	05/01/89 06/06/89 04/27/90	05/01/89 06/06/89	05/02/89	05/02/89
Ship- ment	888	222	57 57 65	57	888	8 8 8	888	8 8	22	22	22.50	27	7.	22
AVS	5119 5119 5119	5121 5121 5121	5138 5138 5138	5142	22 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2	\$176 \$176 \$176	\$197 \$197 \$197	5198 5198	5379 5379	5383 5383	5384 5384 5384	5385 5385	5393	5457 5457

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	Assay	<u> </u>	TTM		H	T.	1		L L	TIM	MIT	I	F	TH	H	H	TTH	I	117	į	TTH	H	MTT	Ę	TTM	HT	H	TTM	H	TI	Ē	MTT	HTT	F		Ē	ĒĒ
	i	ĭ	15.90	8.00	4.97	7 03	27.0	74.0	5.26	4.46	0.05	9	0.0	20.90	5.21	2.70	2.25	0.00	15 20	3	8.50	2.55	42	000	0.21	39.55	9.8	2.37	36,18	0.00	0.61	0.70	35.52	0.74	•	.00	0.00
	;	7	3.58 >	5	0.0	1.05	8 6		8.0	0.20		0	0.0	0.00	0.00	0.00	0.00	0.00	78 0	3	2.14	0.97 >	0 07 >	000	9.0	3.31 >	0.00	0.00	6.02 >	00.0	0.00	0.00	8.48 >	0.0		. 80.0	2.35
	3	\$ \$	0.0	3.5	0.0	00.0	8 6	3	0.00	0.00	0.00	0	0.0	0.00	0.0	0.0	0.00	0.00	2	3	0.00	0.00	5	00.0	0.0	0.0	0.0	0.00	0.00	0.00	0.00	0.00	0.00	0.00	8	88	0.0
	1	5 8	32.00	320.00	100.00	420 00	1000	00.00	3200.00	89.30	32.00	100	100.00	320.00	1000.00	1000.00	320.00	32.00	200 00	20.03	320.00	320.00	200 00	293.00	320.00	32.00	32.00	100.00	320.00	320.00	320.00	320.00	320.00	1000.00	00 002	908.00	320.00 1000.00
		5 8	0.00		0.00	A		•	0.00	0.00	00.00		0.0	٨	٨	^	0.00	0.00	8	3	0.00	0.00	8	00.0	0.00	0.00		0.00	0.00	0.00	0.00	0.00	^ 00.0	0.00		88	0.00
		2 2	4.05	8.8	8	5	3	3	9.0	0.35	0.00	0	0.0	0.0	0.0	0.00	0.00	0.00			3.63	1.33	. 22	0.00	0.00	157.45	0.0	0.00	111.00	0.00	0.0	0.00	8.48	0.0		. 6.	3.67
		16 50	32.00 >	3.4	17.30	320.00 >	973	20.00	2200.00	8.76	32.00	62.20	8.99	320.00 >	00.0001	651.00	320.00	30.60	200 00	250.00	291.00	320.00 >	24.50	8, 70	320.00	32.00 >	32.00	28.20	320.00 >	164.00	93.70	320.00	320.00 >	598.00	730 002	190.00	320.00 734.00
		S 2	7.90 >	**	0.0				A 00.0	24.90	0.00	00	0.00				0.00	0.00	7 84		80.20	240.00 >	100	0.00	0.00	0.20	0.00	0.0	2.88 >	0.00	0.00	0.00	37.70 >	0.00	700 702	9.0	0.00 >
		S IA	5.63	7.7	2.11	1.87	0		20.	3.17	0.00	0.33	0.29	14.10	1.49	0.0	0.00	0.00	1 20	?	3.39	1.57	1 40	0.42	0.00	6.72	0.0	1.26	10.20	0.00	0.00	0.0	37.01	0.00	72.0	0.28	3.32
	1	\$ 21	28.30	8.2	6.74		00 EEY	20.00	1390.0				27.40	320.00 >	735.00	433.00		19.20	70	•	172.00	234.00	7 21	42.20	۳,			17.60	17.30	73.20	54.10	320.00	320,00 >	229.00	5	73.50	100.00
		9	5.03	2.6	3.20	171.00 >	710 00	3 3	3.00	1.55	0.00	2	93.10	22.70 >	767	0.0	0.00	0.00	7 00		50.70	149.00	7 84	100.00	0.00	0.10	0.00	13.90	1.70	0.00	0.0	0.0	8.65	0.0	2	260.00	320.00
			1.238	3.5	0.995	1.268	00	14.0	1.059	0.957	1.553	1 234	1.050	0.950	1.030	1.418	1.271	1.387	1 148	3	1.293	1.225	1 276	1.275	1.260	1.436	1.355	1.029	0.975	1.347	1.105	0.405	90.	1.347	200	1.99	1.056
	Pit				Š		,		727	F				2	I	VIG	AZA	RZY			R2Z	98	5	R32	905	R33	7	N6Z	970	II	28	9	R71	Z		35	AZX
		9160	12/15/89		02/20/90	05/11/90	06/90/24	24/00/20	06/60/	04/17/89	12/15/89	05/11/90	07/20/90	04/17/89	01/05/90	05/04/90	05/11/90	08/11/89	08/21/70	00) IE/01	08/11/89	06/12/89	04/14/80	08/11/89	06/16/89	08/11/89	05/04/90	10/05/90	06/26/89	05/04/90	10/05/90	02/01/89	08/18/89	05/04/90	02/47/00	08/18/89	07/17/89
	Ship-		53	8	*		3 3		8	53	53	3	8	53	53	65	8	20	73	2	99	26	3	26.	26	26	65	26	27	65	57	22	27	65	2	27	57
	AVS		5485		2485		0875		2480		2495		2495			2497		5528						5541			2543			6095		5643		2643		26%	5691

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Acces	179	EEEE	T I	EE	EE	FE	EE	EE	EEE	EE	EE	EE	EE			T I	
	TAI	20.06 20.06 0.83	0.00	6.22	7.83	34.69	2.91	14.84	7.49 9.32 9.94	16.40	39.49	0.00	0.00	11.06	11.23 5.68 12.72	7.92	
	15	0.08 0.08 0.08 0.08	0.00	0.00	9.00	3.62	0.00	3.20 >	1.55 × 0.86 × 1.67	3.70	0.00	1.12 > 0.00	00.0	4.00.0 0.00.0	2.18 1.56 2.44 >	1.30	
	AI 95	8888	0.00	0.0	0.0	22.00	0.00	0.00	0.00	0.00	0.00	80.0	0.0	9000	0.00	0.0	
	70 95	8353	320.00	320.00	31.50	30.50	320.00	320.00 100.00	320.00 32.00 32.00	320.00	320.00	320.00 918.00	259.00	320.00 948.00 838.00	31.00 31.30	247.00	
	26 21	8888	AA	0.00	0.00	9.42	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00 0.00 v	0000	0.00	
	AI 50	0.00 4.28 0.00	0.00	5.12 0.00	0.00	11.95	0.00	57.50 0.00	8.43 11.61 4.69	3.70	0.00	1.62	0.00	0.00	3.31	2.42	
	TC 50	5.67 6.22 6.38 6.18	320.00	320.00 >	19.20	13.10	75.60	57.50 × 25.70	8.43 × 32.00 × 23.30	320.00 >	320.00	320.00 > 183.00	31.00	320.00 × 241.00 156.00	19.20 20.40 21.10	50.90	
	10 50	0.00 2.54 1.49 0.00	A A	62.60 > 0.00	0.00	5.52	0.00	0.00	1.00	86.50 >	20.90	198.00 >	32.00	72.80 v 0.00 0.00	5.79 6.38	18.40	
	VI 25	92.50	0.00	9.6	2.77	11.93	0.17	3.20 < 0.32	1.81	6.78	0.00	1.85	0.54	1.69	3.22 2.65 3.55	2.62	
	10 25	38. 23. 23. 24. 24. 24. 24. 24. 24. 24. 24. 24. 24	126.00	1.00 4	12.40	6.60	32.00	3.20 >	1.55 × 2.36 8.30	320.00 > 447.00	257.00	221.00 71.20	2.79	77.30 66.00 57.50	12.60 14.40 15.60	23.90	
	10 25	0.00	71.00	24.60 < 0.00	4.50	4.11	192.00	1.00	1.31	47.20 >	13.20 >	120.00	5.12	45.60 57.30 61.10	3.92 5.42 4.38	9.09	
	Diff.	0.916 1.007 0.988 1.105	1.158	1.140	0.897	0.897	0.953	1.014 <	1.014 < 1.152 0.975	1.123	1.506	1.170	1.298	1.298 1.421 0.967	1.124 1.385 0.967	1.110	
	Date #	07/31/89 Q22 10/02/89 RV9 10/09/89 RYA 10/05/90 290		10/20/89 S45 02/02/90 TPF	11/03/89 SBO 02/02/90 TNE	11/03/89 SBO 02/02/90 TNF	11/03/89 SBP 02/02/90 TNF	11/10/89 SFV 11/02/90 ZWO	11/10/89 SFV 02/02/90 TP4 02/09/90 TMN	11/10/89 SHX 02/02/90 TP5	01/19/90 TCV 12/07/90 10K	02/16/90 U37 03/23/90 U0U	02/16/90 U38 03/23/90 U0V	02/16/90 U38 03/23/90 U0V 10/05/90 29P	02/16/90 U3A 03/23/90 U0W 10/05/90 29P	02/16/90 U3D 03/16/90 UM3	
Shin-	E)	8888	88	88	22	25	22	22	222	10	62 0	22	22	222	222	63	
	9	5714 5714 5715	325	5854	5905	2009	2065	5997	5998 5998 5998	6059	6218 6218	6309	6310	6311 6313 6311	6315 6315 6315	6321	

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Assay Type	FF	ĒĒ	EE		ĒĒ	EE	EE	EEE	ĒĒ	EE	ĒĒ	ĒĒ	ĒĒ	EE	ĒĒ	EE	EE	
IĀĪ	0.08	3.06	2.91	12.59	9.62	12.30	17.29	3.7	0.00	9.19	0.02	15.32 12.65	16.44	0.57	6.48	1.78	7.11	
3	0.26 >	0.00	0.56	2.29	16.26 > 0.91 >	27.38 > 0.00	0.00	0.00	0.27 >	0.00 \$	0.00	1.85	2.40 > 1.14 >	0.00	1.07 >	0.74 >	1.58 > 0.00	
A1 95	0.0	0.00	0.0	0.00	0.00	0.00	3.39	988	0.0	0.00	0.00	0.00	0.00	0.00	0.00	0.0	0.00	
70 %	320.00 973.00	316.00 320.00	274.00	308.00	31.20	320.00 320.00	320.00 >	320.00 320.00 100.00	320.00 320.00	320.00	320.00	318.00 320.00	98.10 95.80	320.00 932.00	320.00 970.00	320.00 938.00	320.00	
20 20	0.00	0.00	0.00	0.0	0.00	0.00	%.40 ^ 0.00 ^	0.00	0.00	0.00	0.00	0.0	0.00	0.00	0.00	0.00	0.00	
Ai 50	0.00	1.71	2.33	3.32	24.77	39.57	0.00	9.85 0.00 0.00	1.28	0.00	0.00	2.53	3.54	3.97	1.54	1.08	0.00	
1C 50	308.00	157.00	57.90	198.00 71.50	66.90	320.00 >	0.93	83.00 29.80 79.60	320.00 > 91.10	320.00	76.20	215.00	60.90	320.00	320.00 >	320.00 >	320.00 >	
10 50	278.00	91.60	24.90	59.60	2.9	0.00	55.90	8.43 0.00 0.00	249.00 > 0.00	0.00 >	0.00	85.00	17.20	0.00	208.00 > 227.00	296.00 >	203.00 >	
AI 25	0.42	0.00	1.23	3.85	27.60	00.0	0.00	1.21	0.77	0.00	0.52	7.39	6.70	0.00	2.53	1.43	2.28	
70 25	100.00	82.00	14.00	137.00	43.90	221.00	0.46	5.01 5.27 8.00	68.20 46.30	> 320.00 > 1000.0 >	47.90	157.00	41.40	230.00	222.00	219.00	> 320.00 > 353.00	
16 25	156.00	53.00	11.30	35.50	1.59	0.00	41.80	4.14 5.59 0.00	88.50 320.00	0.00	91.90	21.30	6.18	0.00	87.70 129.00	153.00	140.00	
Diff.	1.122	1.323	1.182	1.200	1.245	0.999	0.912	1.020	1.158	1.022	1.169	1.151	0.863	1.058	1.442	1.342	1.394	
=	0 UPT	0 W25	0 KEU	N WTE	0 W71	O LYK	0 KA2	0 MJS 00 XG0 00 170	0 UGL 0 VU3	O XGU	90	NAX O	O XN3	0 XQY	O WZR	0 WZU 00 XZ1	0 WZV	
Test	03/02/90 UBT 03/23/90 UP1	05/11/90	05/18/90 07/13/90	05/18/90	05/18/90 U7I 07/13/90 XEK	05/25/90 U9K 07/13/90 XEP	05/25/90 V90 07/06/90 XAJ	06/08/90 07/13/90 11/16/90	03/09/90 UGL 05/04/90 VU3	06/08/90 MM0 07/13/90 XGU	06/15/90 Wa6 07/13/90 xGV	06/15/90 WOH 08/03/90 XYM	09/119/90	07/27/90	06/22/90	06/22/90	06/22/90 WZV 08/10/90 XZ2	
Ship-	33	33	88	88	88	88	88	79 79 79	88	67	67	33	69	66	33	22	88	
AVS.	£53 £73	54.12	47	64.82	64.89	6521 6521	6532	6724 6724 6724	6739	8829	1679	6837	9069	6943	9869	6993	7669	

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Assay Type		EE	EE	FF	TH	EE	ĒĒ	EEE			EEE	EEE			E E
Ţ.	9.19	7.7	10.35	12.26	9.14	10.60	23.46	2.53 0.35 1.00	2.04 4.52 9.89	1.11	383	0.00	16.51	0.00	0.00
13	1.69 >	0.88 >	1.62 >	0.00	1.17 > 0.00	0.00	3.20 >	0.00	0.00 > 2.06 >	0.00 0.00 3.93 v	993	0.00	10.90 •	0.00	1.08 >
A1 %	8.8	9.0	0.00	0.00	8.00	9.0	88	988	0.00	0.00	989	9.00	0.00	0.00	0.00
70 %	320.00	320.00	320.00	320.00 320.00	320.00	320.00 308.00	309.00	320.00 1000.00 1000.00	320.00 314.00 320.00	100.00 320.00 100.00 *	320.00 100.00 100.00	93.30 100.00 100.00	1000.00	1000.00	1000.00
25	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00 .00.00 .00.00	0.00	0.00	0.00	0.00	0.00
AI 50	1.69	1.19	0.00	6.16	1.17	1.13	0.00	0.36	0.00	0.00 2.98 3.93	0.00	0.00	0.00	0.00	1.08
TC 50	320.00 >	320.00 >	320.00 × 575.00	320.00 × 178.00	320.00 >	320.00 > 200.00	210.00	64.00 1000.00 1000.00	320.00 168.00 320.00 >	100.00 177.00 100.00 •	130.00 15.70 100.00 >	33.30 43.30 100.00 >	1000.00	1000.00	1000.00
10 50	190.00	269.00 > 216.00	197.00 > 0.00	76.90 v 0.00	273.00 \$	283.00 > 0.00	00.0	0.00 0.00 0.00	0.00 × 73.80 × 93.50 ×	0.00 > 59.30 > 25.50 >	20.80	20.00	0.00 >	2100.00 >	2960.00 >
AI 25	2.57	1.82	2.54	0.00	2.57	3.76	15.50	5.51 0.00 0.00	0.00 2.06 3.59	0.00	0.00	0.24 0.65 3.54	1.61	0.00	0.00
25 27	320.00	237.00	320.00 >	220.00 31.20	320.00 >	253.00	155.00 155.00	320.00 690.00 1000.0	320.00 80.60 192.00	100.00 23.80 100.00 *	68.30 6.78 13.90	6.78 11.80 40.00	1000.0 >	200.00	2.76
10.25	125.00	130.00	126.00 > 245.00	00.00	125.00 >	67.30	0.00	58.10 0.00 0.00 v	0.00 × 39.20 × 53.50	0.00 × 21.20 15.10 ×	0.00 24.90 6.04	28.60 18.10 11.30	622.00 >	0.00	0.00
Diff.	1.107	1.003	1.003	0.849	0.849	1.092	1.092	1.2%	1.263 1.103 1.152	1.275	1.275	1.254 1.127 1.074	1.038	1.138	1.293
==	XTC YES	XTY 7L6	XTY YLD	1. K	X X X X X X X X X X X X X X X X X X X	XY.	XX	YDA 27.5 29.7	7FC 27N 27N 2PC	222	YFD 270	YFE 27P 2PE	Y12 YS2	78F 220	79R 22T
Test Date	07/27/90 XTU 09/07/90 YLB	07/27/90	04/27/90	08/03/90	06/03/60	08/03/90	08/03/90	08/28/90 10/05/90 10/26/90	08/28/90 10/05/90 10/26/90	08/28/90 10/05/90 10/26/90	08/28/90 10/05/90 10/26/90	08/28/90 10/05/90 10/26/90	08/10/90	08/24/90	08/24/90
Ship-	\$\$	\$ \$	69	69	69	69	69	222	222	222	222	222	22	22	22
AVS S	6002	9 9107	6 7107 6 7107	7031 6	7032 6	7047 6	7048 6	7059 7 7059 7 7059 7	7083 7083 7083	7085 7085 7085	7086 7086 7086 7	7087 7087 7087	7 3117	7386 7	7403 7
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Assay	H	EE	==	ËË	==
¥	6.53	2.29	1.43	2.74	9.72
2	0.00 \	6.3	0.00 > 0.27 >	0.00	0.00
A1 95	88.	8.00	88	0.0	0.00
10 95	1000.00	32.00	320.00	0.00 > 3200.00	1930.00
S 21	0.00	0.00	0.00	0.00	0.00
AI 50	0.00	0.00	1.8	1.12	1.36
TC 50	554.00	32.00 >	3200.00	3200.00 >	1260.00
10 50	2150.00 >	0.00	0.00 > 320.00 > 2390.00 >	2870.00 > 3200.00 > 0.00 1060.00	2390.00 > 3
AI 25	2.88	0.00	0.00	0.16	3137
22 21	262.00 3200.0	4.90	320.00	265.00	891.00 3200.0 >
10 25	269.00	0.00	736.00	1660.00	950.00
Diff.	1.054	1.041	1.158	1.056	1.035
=	ZPH	THE SYS	25F 22P	17E	120
Test	09/21/90 10/26/90	09/21/90	10/12/90	11/16/90 17E 12/04/90 1H1	11/16/90
Ship- ment	KK	RR	22	ĸĸ	KK
AVS.	33	2,60	21.6	1767	8212 8212

The differential is the difference in the cell control and the virus control optical densities. DIFRHTL =

1C25,50,95

(Viral) inhibitory concentration 25%, 50% and 95% = The drug concentration (kg/ml) that inhibited viral CPE by 25%, 50% or 95% calculated by using a regression analysis for semilog curve fitting.

(Cell) toxicity concentration 25%, 50% and 95% = The drug concentration (µg/ml) that reduced cell viability by 25%, 50% or 95% TC25,50.95 Antiviral Index = A single point ratio of the antiviral and anticellular effect of the compound, calculated with 25%, 50% or 95% reduction values (calculated by dividing the $TC_{25,50,95}$ by the $TC_{25,50,95}$). AI 25.50.95

Selectivity Index = A ratio calculated by dividing the TC₂₅ by the IC₅₀ (based upon 6 one-half-log₁₀ dilutions, µg/ml, the maximum scale is 0-320).

Total Antiviral Index = The area between the cytotoxicity and the antiviral curves (based upon a scale of 0-100X).

TAI

S

Activity = A "+" denotes a rest that produced >25% reduction in CPE. A "-" denotes an inactive test (i.e. <25% reduction in CPE).

4.1.6 Punta Toro Virus (PT):

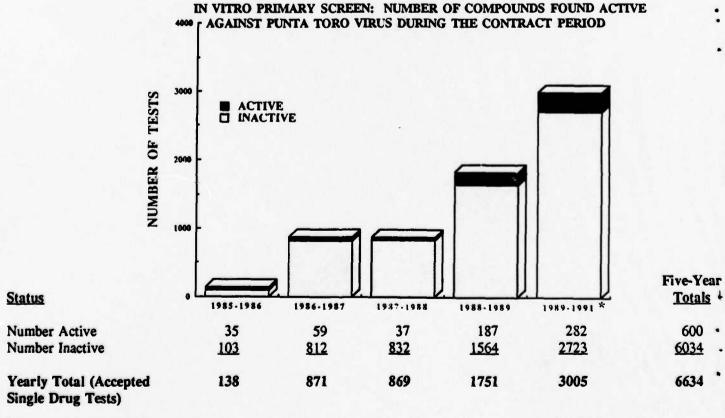
The number of single drug tests carried out against PT during this contract period is summarized in yearly increments in Figure 30. During this five-year period two main in vitro antiviral assay protocols were implemented:

- 1. A standard CPE inhibition assay by virus rating (VR) (Annual Report, December 15, 1988, Section 3.2.4).
- 2. Since November, 1988, MTT based-antiviral assay format.

A total of 8221 tests were performed during this contract period using both assay types. Routine testing was changed to the MTT-assay format to improve the efficiency and quality of the primary screening program in addition to being more cost-effective. Ribavirin (AVS-0001) was tested in each standard virus rating (VR) CPE-inhibition assay as a positive control compound. Results of these positive control (VR tests) were used as a guideline to assess the quality of each assay.

After the testing was converted to the MTT-assay format, we performed a total of 406 control compound assays with Ribavirin during the last 26 months of the contract period. During this time 684 tests were internal (+++) virus load, cell load, and other quality control tests. Four hundred thirty-eight (438) tests were considered unsatisfactory based on the criteria of the quality controls set during this reporting period. The rest, totaling 4756 were actual single drug MTT-assays. The total number of MTT-assays (6284) tested during the last two years represents a 224% increase (improvement) in the total testing output as compared to the total of 1937 tests performed during the first 3 years of this contract.

Out of the 6693 accepted single drug tests, 635 compounds demonstrated antiviral activity at greater than 50% reduction levels. This represents around 9.5% of the tested compounds having *in vitro* antiviral activity against PT-virus. The remainder, 6058 compounds (90.5%), were considered inactive with both assay protocols (Figure 30).



Represents 14-month period (November 15, 1989 - January 31, 1991)
Figure 30

- 4.1.6.1 <u>PT-Quality Controls (MTT Assay):</u> Two positive control compounds (Ribavirin and 2-Thio-6 Azauridine) were used in the daily assay sets as antiviral activity quality controls. The antiviral performance of the unknown compounds is compared to that of the positive control compounds. Compounds with equal to of better antiviral potency are considered active and are worthy of further in vitro profile studies and in vivo testing.
- 4.1.6.1.1 Antiviral Activity of Ribavirin vs PT Virus: A summary of the antiviral and cytotoxicity performance of the primary control compound, AVS-0001 (Ribavirin) is presented in Figure 31-A for 242 tests performed during November, 1989 through January, 1991.

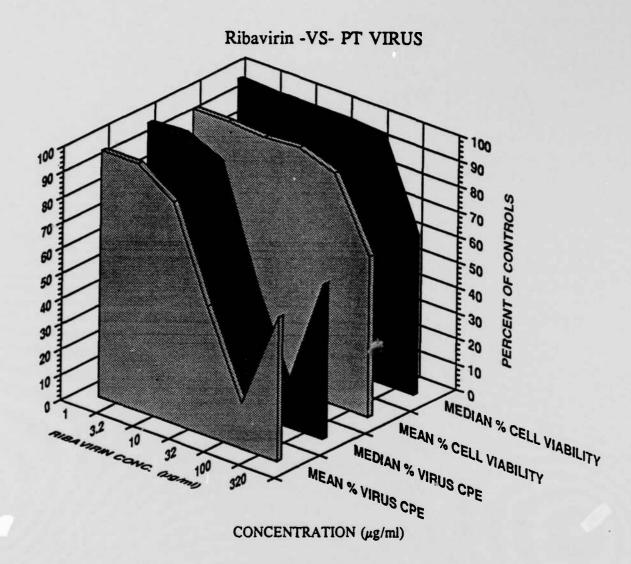
<u>Control Compound-Antiviral Performance</u>: Ribavirin (AVS-0001) has been the sole control compound against PT in these MTT-assay screens. The mean and median antiviral inhibition and cytotoxicity patterns of the positive control drug (Ribavirin) are illustrated in Figure 31-A.

The 242 control tests performed with Ribavirin gave a mean Total Antiviral Index (TAI) of 27.37% (SD \pm 12.56) and the median value was 26.49%. The TAI measures the overall antiviral effectiveness of the compound and it ranged from -(3-70)% during this period. The mean Selectivity Index (SI) was only 8.81 (SD \pm 8.33) and the median SI value was 6.48, indicating moderate antiviral selectivity for Ribavirin against PT virus. The SI ranged from -(0.01-57) during this period. However, the closeness of the mean and median values indicate that the present execution of the SOP is consistent and repeatable.

The mean Antiviral Index 25% (AI₂₅) value was 19.39 (SD \pm 31.54). The median AI₂₅ value was 12.36 (range 0.02 - 320). The mean Antiviral Index 50% (AI₅₀) was 12.07 (SD \pm 10.14) with a median of 8.76 (range 1.65 - 65.61). The mean Antiviral Index 95% (AI₉₅) was 1.87 (SD \pm 2.98), with a median of 0 (range 0 - 12.74). This indicates that the control compound, Ribavirin, does not consistently reach 95% antiviral reduction levels.

The mean Antiviral Inhibitory Concentration 25% (IC₂₅) was 21.02 μ g/ml (SD \pm 13.40). The median IC₂₅ value was 16.80 μ g/ml (range = 1 - 72.9 μ g/ml). The mean Antiviral Inhibitory Concentration 50% (IC₅₀) was 38.47 μ g/ml (SD \pm 24.69). The median IC₅₀ value was 33.30 μ g/ml (range = 3.0 - 194 μ g/ml). The mean Antiviral Inhibitory Concentration 95% (IC₉₅) was 34.15 μ g/ml (SD \pm 50.26). The median IC₉₅ value was 0 μ g/ml (range from 0 - 320 μ g/ml). This discrepancy indicates that the control compound Ribavirin does not consistently reach 95% reduction levels. During this reporting period, the highest starting concentration of Ribavirin (100 μ g/ml)was increased from the previous high dose of 100 to 320 μ g/ml to properly evaluate the maximum antiviral effect of Ribavirin.

The average maximum antiviral inhibitory level of 242 Ribavirin tests (Figure 31-A) was reached at 100 μ g/ml of the compound with 85% antiviral effect. Maximum antiviral effect (~89%) was found with a simultaneous ~5% cytotoxic suppression. Above (100 μ g/ml) concentration Ribavirin starts to lose its antiviral potency (~40%) at 320 μ g/ml while simultaneously the Ribavirin becomes maximally toxic (~40%).



76 VII al CI L	70 Cell Viability
% Viral CPE	% Cell Viability

Conc.(µg/ml)	1	3.2	10	32	100	320	1	3.2	10	32	100	320
Mean	97	97	87	51	18	57	96	96	95	96	91	63
Median	99	100	94	51	16	61	99	98	98	98	97	62
Std. Dev.	0.05	0.07	0.17	0.28	0.18	0.26	0.06	0.05	0.05	0.05	0.12	0.19

Figure 31-A

Average Antiviral and Cytotoxicity Values for 242 Positive Control Compound Tests

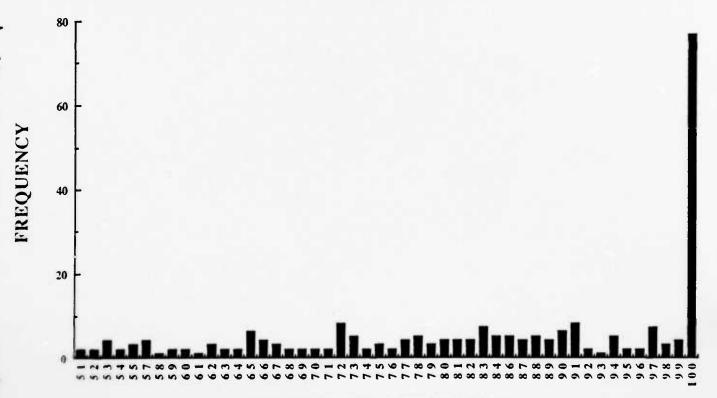
4.1.6.1.3 <u>Maximum Antiviral Effect of Ribavirin vs PT Virus:</u> Since the metabolic activity of the cells was an unknown function during the testing period, it was monitored indirectly by measuring the maximum antiviral effect of the control compound Ribavirin. This demonstrated the amount of infectious virus that was produced by the cells (Maximum Percent CPE).

A bar graph scatter plot (Figure 32-A) depicts the distribution of the maximum antiviral reduction values of all 242 control compound assays for Ribavirin. The results indicate that the average maximum antiviral reduction obtained with the present SOP is around 85% (SD \pm 15.14) reduction levels. The maximum reduction levels vary from 51 - 100% but remain quite consistently around the median of 89%. The assay control values give a shifted half-bell-shaped distribution curve toward the maximum 100% reduction level. This indicates quite a consistent day-to-day performance of the control compound in the PT-MTT assay.

During this period the positive control compound performance criteria for Ribavirin versus the PT virus was set at 50% reduction level. All assays in which Ribavirin did not meet this accepted quality control level (\geq 50%) were rejected (i.e., 438 unsatisfactory tests).

Ribavirin is active *in vitro* against PT virus and functions as a reasonable quality control compound. On the other hand, regardless of the performance of the PT-quality control drug Ribavirin, around 282 other compounds have equal or better antiviral activity against PT virus than AVS-0001. (See 95% and 50% reduction summaries).

VARIATION OF THE MAXIMUM ANTIVIRAL EFFECT PT VIRUS - VS - Ribavirin



PERCENT CPE REDUCTION

Figure 32-A

Maximum Antiviral CPE Reduction (%).

Summary of 242 Control Tests.

4.1.6.1.4 Cellular Cytotoxicity of Ribavirin vs PT Virus:

<u>PT-Control Compound-Cytotoxicity Performance:</u> The 242 cytotoxicity values of the positive control compound Ribavirin are also very consistent. The mean cell Toxic Concentration 25% (TC₂₅) was 225.78 μ g/ml (SD \pm 87.67) and the median was 241 μ g/ml (range of 0.7 - 320 μ g/ml). The mean cell Toxic Concentration 50% (TC₅₀) value was 303.7 (SD \pm 35.76) and the median was 320 μ g/ml (range of 119-320 μ g/ml). The mean cell Toxic Concentration 95% (TC₅₅) value was 320 (SD \pm 1.1) and the median was 320 (range of 307-320 μ g/ml). This discrepancy indicates that the control compound Ribavirin does not consistently reach 95% cytotoxicity levels.

As can be seen from Figure 31-A, the toxicity starts to become measurable above the concentration of 100 μ g/ml and the maximum toxicity has not been reached at 320 μ g/ml. Further increase of the concentration of Ribavirin would be needed to properly evaluate the maximum cytotoxicity of Ribavirin.

Also Figure 31-A, indicates that when the cytotoxicity reaches ~ 0 - 5% at 100 μ g/ml, the control compound (Ribavirin) has reached simultaneously its maximum antiviral effect (85%). The cytotoxic effect of Ribavirin is insignificant between 1 and 100 μ g/ml. The maximum cytotoxicity reached $\sim 40\%$ at 320 μ g/ml, which is the highest Ribavirin concentration tested.

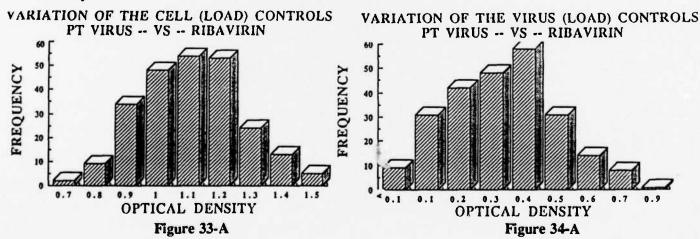
Ribavirin has a definite cytotoxic suppression on cellular metabolism and growth. However, the TC_{25} and TC_{50} toxicity could not be consistently achieved with the 100 μ g/ml concentration of Ribavirin. Therefore, a readjustment to 320 μ g/ml (as being the highest Ribavirin concentration tested) was done during this reporting period. However, at this concentration (320 μ g/ml) the TC_{50} and TC_{95} cannot yet be measured consistently.

4.1.6.1.5 PT-Assay Plate Quality Controls: Cell Load and Virus Load Parameters (Ribavirin): The MTT assay is fundamentally dependent upon the quality of the assay plates. Our large-scale antiviral testing is dependent upon the uniformity of the test plates produced for the daily assays. Equal numbers of cell load and virus load as well as the consistent performance of the reagents used daily was monitored. A sample of the plate variation control for the period of November, 1989 through January, 1991, is presented in Figures 33-A, 34-A and 35-A.

PT-Control Compound-Cell Load Performance: A bar graph scatter plot of the mean cell control (O.D. reading) of 242 control assays is plotted in Figure 33-A. The results indicate that the cell O.D. readings reached a mean of 1.110 (SD \pm 0.160) with a median of 1.110 (range of 0.720 - 1.540). This indicates that a uniform and equal number (18,000 cells/well) of cells are being loaded into every well in the 96-well plate during the day-to-day operation. The cells reduced MTT to formazan giving maximum blue color uniformly and consistently.

<u>PT-Control Compound-Virus Load Performance</u>: A bar graph scatter plot of the mean virus load O.D. readings of the 242 control assays is presented in Figure 34-A. The results indicate that the average load O.D. reading is 0.330 (SD \pm 0.170) with a median of 0.330 (range of 0.004 - 0.920). This demonstrates that a good cell destruction is taking place and a uniform load of virus (32 TCID₅₀) is administered on the cell monolayer with very consistent viral CPE results.

PT-Control Compound-Assay Differential Performance: A bar graph scatter plot of the mean O.D. differential values of the 242 control assays is provided in Figure 35-A. The results indicate that the average differential O.D. reading is 0.777 (SD \pm 0.190) with a median of 0.774 (range 0.327 - 1.500). The single bell-shaped curve is reasonably sharp and uniform. This reflects that the assays are executed consistently and are repeatable during day-to-day operation with close to 78% measurement accuracy.



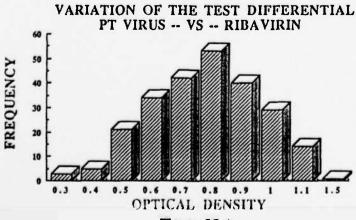


Figure 35-A

4.1.6.1.1 Antiviral Activity of AVS-6724 (2-Thio-6-Azauridine) vs PT Virus: A summary of the antiviral and cytotoxicity performance of the second control compound, AVS-6724 (2-Thio-6-Azauridine) is presented in Figure 31-B for 44 tests performed during November, 1989 through January, 1991.

Second Control Compound-Antiviral Performance: 2-Thio-6-Azauridine (AVS-6724) has been tested as a possible second control compound against PT in these MTT-assay screens. The mean and median antiviral inhibition and cytotoxicity patterns of this second positive control drug are illustrated in Figure 31-B.

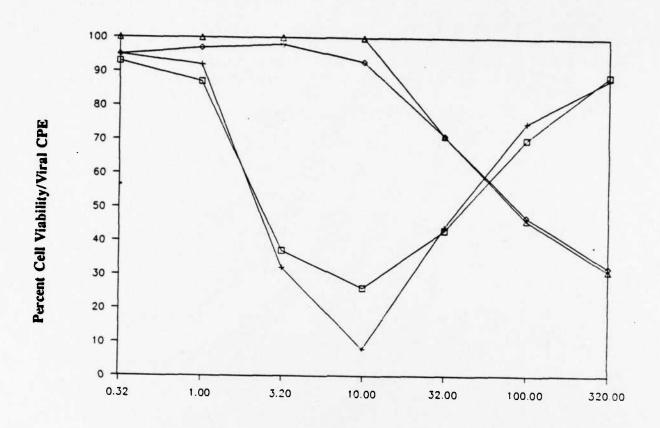
The 44 control tests performed with 2-Thio-6-Azauridine gave a mean Total Antiviral Index (TAI) of 36.00% (SD \pm 15.20) and the median value was 34.90%. The TAI measures the overall antiviral effectiveness of the compound and it ranged from \sim 11.54 - 69.30% during this period. The mean Selectivity Index (SI) was only 15.69 (SD \pm 13.50) and the median SI value was 12.32, indicating moderate antiviral selectivity for 2-Thio-6-Azauridine against PT virus. The SI ranged from \sim 0 - 63.65 during this period. However, the closeness of the mean and median values indicate that the present execution of the SOP is consistent and repeatable.

The mean Antiviral Index 25% (AI₂₅) value was 25.10 (SD \pm 20.60). The median AI₂₅ value was 18.70 (range 5.02 - 92.14). The mean Antiviral Index 50% (AI₅₀) was 37.10 (SD \pm 33.90) with a median of 33.40 (range 0 - 172.79). The mean Antiviral Index 95% (AI₉₅) was 16.80 (SD \pm 24.70), with a median of 10.70 (range 0 - 107.68). This indicates that the, 2-Thio-6-Azauridine, does not consistently reach 95% antiviral reduction levels.

The mean Antiviral Inhibitory Concentration 25% (IC₂₅) was 1.85 μ g/ml (SD \pm 1.20). The median IC₂₅ value was 1.40 μ g/ml (range = 0.33 - 5.81 μ g/ml). The mean Antiviral Inhibitory Concentration 50% (IC₅₀) was 3.10 μ g/ml (SD \pm 2.30). The median IC₅₀ value was 2.20 μ g/ml (range = 0 - 12.4 μ g/ml). The mean Antiviral Inhibitory Concentration 95% (IC₉₅) was 3.65 μ g/ml (SD \pm 3.90). The median IC₉₅ value was 3.0 μ g/ml (range from 0 - 9.6 μ g/ml). This discrepancy indicates that the 2-Thio-6-Azauridine does not consistently reach 95% reduction levels. During this reporting period, the highest starting concentration of 2-Thio-6-Azauridine (320 μ g/ml) was varied to properly evaluate the maximum antiviral effect of 2-Thio-6-Azauridine. The best window appears to be from a high concentration of 100 μ g/ml to a low concentration of 0.320 μ g/ml in six 1/2 log₁₀ increments. See Figure 31-B. At this scale (0.32 - 100 μ g/ml) all important antiviral (IC₂₅, IC₅₀, IC₉₅ and TAI) parameters are measured as well as all important cytotoxicity parameters (TC₂₅, TC₅₀, except TC₉₅) as also indicated.

The average maximum antiviral inhibitory level of 44, 2-Thio-6-Azauridine, tests (Figure 31-B) was reached at $10 \mu g/ml$ of the compound with 90% antiviral effect. Maximum antiviral effect (~98%) was found with a simultaneous ~10% cytotoxic suppression. Above $10 \mu g/ml$ concentration 2-Thio-6-Azauridine starts to lose its antiviral potency (~10%) at 320 $\mu g/ml$, while simultaneously the 2-Thio-6-Azauridine becomes maximally toxic (~70%) with increasing cytotoxicity.

2-THIO-6-AZAURIDINE -VS- PT VIRUS



CONCENTRATION (µg/ml)

☐ Mean %	+ Median %	♦ Mean % Cell	ΔMedian % Cell
Viral CPE	Viral CPE	Viability	Viability
	% Viral CPE		% Cell Viability

Conc.(µg/ml)	0.32	1	3.2	10	32	100	320	0.32	1	3.2	10	32	100	320
Mean	93	87	37	26	43	70	89	95	97	98	93	71	47	32
Median	95	92	32	8	44	75	88	100	100	100	100	71	46	31
Std. Dev.	0.07	0.19	0.32	0.19	0.22	0.18	0.10	.07	.06	0.05	0.11	0.17	0.11	0.04

Figure 31-B
Average Antiviral and Cytotoxicity Values for 44 Positive Control Compound Tests

4.1.6.1.3 <u>Maximum Antiviral Effect of 2-Thio-6-Azauridine vs PT Virus:</u> Since the metabolic activity of the cells was an unknown function during the testing period, it was monitored indirectly by measuring the maximum antiviral effect of the control compound 2-Thio-6-Azauridine. This demonstrated the amount of infectious virus produced by the cells (Maximum Percent CPE).

A bar graph scatter plot (Figure 32-B) depicts the distribution of the maximum antiviral reduction values of all 44 control compound assays for 2-Thio-6-Azauridine. The results indicate that the average maximum antiviral reduction obtained with the present SOP is around 88% (SD \pm 16.40) reduction levels. The maximum reduction levels vary from 47 - 100% but remain quite consistently around the median of 99%. The assay control values give a shifted half-bell-shaped distribution curve toward the maximum 100% reduction level. This indicates quite a consistent day-to-day performance of the control compound in the PT-MTT assay.

Recommendations:

Based upon the data obtained in parallel studies with Ribavirin, we recommend that 2-Thio-6-Azauridine (AVS #6724) will be used as a second control compound against PT virus. It's overall performance is slightly better than the present control, Ribavirin. It is readily available from Sigma Chemical Company, it is inexpensive and works at ~10-fold lower concentrations than Ribavirin.

VARIATION OF THE MAXIMUM ANTIVIRAL EFFECT PT VIRUS - VS - 2-THIO-6-AZAURIDINE

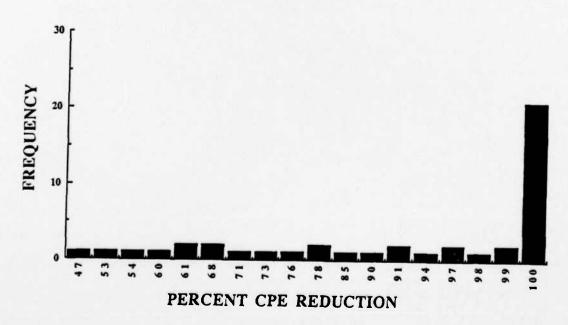


Figure 32-B

Maximum Antiviral CPE Reduction (%).

Summary of 44 Control Tests.

4.1.6.1.4 Cellular Cytotoxicity of 2-Thio-6-Azauridine vs PT Virus:

PT-Control Compound-Cytotoxicity Performance: The 44 cytotoxicity values of the positive control compound 2-Thio-6-Azauridine are also very consistent. The mean cell Toxic Concentration 25% (TC₂₅) was 35.60 μ g/ml (SD \pm 22.9) and the median was 28.7 μ g/ml (range of 7.63 - 87.9 μ g/ml). The mean cell Toxic Concentration 50% (TC₅₀) value was 78.8 (SD \pm 27.4) and the median was 90.1 μ g/ml (range of 22.3 - 120 μ g/ml). The mean cell Toxic Concentration 95% (TC₉₅) value was 155 (SD \pm 96.4) and the median was 100 (range of 100 - 320 μ g/ml). This discrepancy indicates that the control compound 2-Thio-6-Azauridine does not consistently reach 95% cytotoxicity levels.

As can be seen from Figure 31-B, the toxicity starts to become measurable above the concentration of 10 μ g/ml and the maximum toxicity has not been reached at 320 μ g/ml. Further increase of the concentration of 2-Thio-6-Azauridine would be needed to properly evaluate the maximum cytotoxicity of 2-Thio-6-Azauridine.

Also Figure 31-B, indicates that when the cytotoxicity reaches ~ 0 - 10% at $\sim 10 \,\mu\text{g/ml}$. The control compound (2-Thio-6-Azauridine) has reached simultaneously its maximum antiviral effect (90%). The cytotoxic effect of 2-Thio-6-Azauridine is insignificant between 1 and 10 $\mu\text{g/ml}$. The maximum cytotoxicity reached $\sim 70\%$ at 320 $\mu\text{g/ml}$, which is the highest 2-Thio-6-Azauridine concentration tested.

2-Thio-6-Azauridine has a definite cytotoxic suppression on cellular metabolism and growth. However, the TC_{25} and TC_{50} toxicity could be consistently achieved with the 100 μ g/ml concentration of 2-Thio-6-Azauridine. Therefore, an adjustment to 320 μ g/ml as being the highest 2-Thio-6-Azauridine concentration tested.

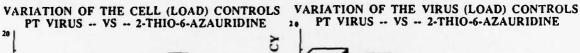
4.1.6.1.5 PT-Assay Plate Quality Controls: Cell Load and Virus Load Parameters (2-Thio-6-

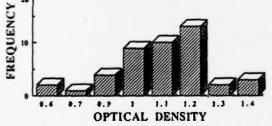
Azauridine: The MTT assay is fundamentally dependent upon the quality of the assay plates. Our large-scale antiviral testing is dependent upon the uniformity of the test plates produced for the daily assays. Equal loads of cell load and virus load as well as the consistent performance of the reagents used daily was monitored. A sample of the plate variation control for the period of November, 1989 through January, 1991 is presented in Figures 33-B, 34-B, and 35-B.

Second PT-Control Compound-Cell Load Performance: A bar graph scatter plot of the mean cell control (O.D. reading) of 44 control assays is plotted in Figure 33-B. The results indicate that the cell O.D. readings reached a mean of 1.100 (SD \pm 0.170) with a median of 1.130 (range of 0.610 - 1.440). This indicates that a uniform and equal number (18,000 cells/well) of cells are being loaded into every well in the 96-well plate during the day-to-day operation. The cells reduced MTT to formazan giving maximum blue color uniformly and consistently.

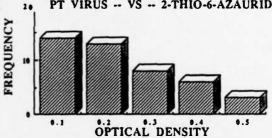
Second PT-Control Compound-Virus Load Performance: A bar graph scatter plot of the mean virus load O.D. readings of the 44 control assays is presented in Figure 34-B. The results indicate that the average load O.D. reading is 0.230 (SD \pm 0.130) with a median of 0.210 (range of 0.060 - 0.540). This demonstrates that a good cell destruction is taking place and a uniform load of virus (32 TCID₅₀) is administered on the cell monolayer with very consistent viral CPE results.

Second PT-Control Compound-Assay Differential Performance: A bar graph scatter plot of the mean O.D. differential values of the 44 control assays is provided in Figure 35-B. The results indicate that the average differential O.D. reading is 0.860 (SD \pm 0.200) with a median of 0.880 (range 0.199 - 1.253). The single bell-shaped curve is reasonably sharp and uniform. This reflects that the assays are executed consistently and are repeatable during day-to-day operation with close to 86% measurement accuracy.





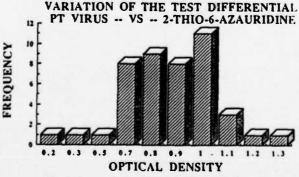
Variation of the Cell (Load) Controls PT Virus -- VS --2-Thio-6-Azauridine



Variation of the Virus (Load) Controls PT Virus - VS - 2-Thio-6-Azauridine

Figure 33-B

Figure 34-B



Variation of the Test Differential PT Virus -- VS -- 2-Thio-6-Azauridine

Figure 35-B

4.1.6.2 PT-Antiviral Activity Results:

New Drugs with 95% Antiviral Reduction Levels: Out of the 6693 actual single drug tests, 107 new compounds demonstrated excellent antiviral activity, having antiviral reduction values of equal to or better than 95%. This represents around 1.6% of the test compounds being active at this excellent reduction level. These compounds are summarized in Table 16 according to the highest Total Antiviral Index (TAI). Compound AVS-5580 demonstrated the greatest in vitro promise, having a TAI of 96% and Selectivity Index (SI) of > 320. The next fourteen compounds, AVS-6229, 2309, 4848, 4855, 5546, 6176, 3802, 6197, 0217, 6219, 8353, 6195, 4863 and 1644, demonstrated excellent antiviral activity with TAI's greater than 50% and SI values that ranged from 19 - 222. Fourteen other compounds demonstrated good antiviral activity, having TAI's from 40 - 49% and SI's from 9 - 41. The rest (78 compounds) had only moderate antiviral activity with TAI's ranging from 10 - 39% and SI's of < 1 to 11.

It is worthwhile to note that compounds received in shipment number 62 were mostly colored (Table 16). Therefore those compounds appearing in the 95% active category from shipment number 62 should be interpreted with caution, since colored compounds create false positive readings with the MTT assay.

Table 16

AVS Compounds Active Against Punta Toro Virus (PT) at AI₉₅ Level

	AVS	Ship-	Test	Diff-										
Virus	No.	ment#	Date	rntl.		IC 9	5	TC 95	;	AI 95	,	SI		TAI
	5580	54	05/03/89			7.26				13800.0				-
	6229	62	01/16/90		<	1.00		320.00			>	221.58		
	2309	53	04/11/89			16.80		320.00		19.10		53.10		
	4848	48	02/22/89	0.429		71.50	>	320.00	>		>	104.00		
	4855	48	02/22/89			16.80		328.00		19.50		20.50	>	
	5546	56	06/14/89			7.96		320.00				151.00		60.74
	6176	62	01/09/90			143.00		320.00		2.23		32.84		
	3802	67	07/12/90			0.84		10.00		11.95		42.71		
	6197	62	01/11/90			44.30		320.00		7.22		21.34		
	0217	61	12/06/89			123.00		320.00		2.61		22.96		
	6219	62	01/16/90			144.00		320.00		2.22		21.52		
	8353	75	11/29/90			31.60		320.00		10.14		19.05		
PT	6195	62	01/11/90			168.00	>	320.00	>	1.90	>	19.85		
PT	4863	48	02/22/89	0.405		27.80		320.00		11.50		129.00	>	50.97
PT	1644	64	05/03/90			79.60	>	1000.00	>	12.56		26.49		50.83
PT	5494	53	04/18/89			29.40	>	100.00	>	3.40		40.90	>	48.61
PT	4871	61	12/06/89	0.800		239.00	>	1000.00	>	4.18	>	22.14	>	48.44
PT	6178	62	01/09/90	0.607		88.40	>	320.00	>	3.62	>	11.81	>	46.18
PT	6202	62	01/11/90	0.763		90.50	>	320.00	>	3.54	>	8.69	>	45.62
PT	5138	57	07/11/89	0.554		89.00	>	320.00	>	3.59	>	10.50	>	44.37
PT	6218	62	01/16/90	0.777		30.10	>	320.00	>	10.63		17.51	>	44.15
PT	5780	59	01/23/90	0.824		71.50		318.00		4.44		17.99		43.57
PT	6175	62	01/09/90	0.636		95.40	>	320.00	>	3.36	>	9.77	>	43.48
PT	5058	48	02/28/89	0.548		320.00	>	320.00	>	1.00	>	12.00	>	42.55
PT	7084	72	10/25/90	0.796		9.31		95.80		10.28		12.88	>	42.40
PT	5067	48	03/01/89	0.886		90.00	>	320.00	>	3.56	>	9.19	>	41.88
PT	3494	33	04/19/89			9.79		96.40		9.85		12.20		41.87
PT	5601	GABSN				846.00	>	3200.00	·>	3.78		14.77	>	40.14
PT	5040	45	12/20/89			83.70	>	1000.00	>	11.95		14.55		39.83
PT	5497	53	04/18/89	0.672		89.80	>	320.00	>	3.56	>	9.38	>	39.16
PT	5053	48	02/28/89	0.452		10.00		285.00		28.50		9.88		39.13
PT	3586	32	04/19/89			29.00	>	320.00	>	11.00		0.27		37.49
PT	5515	53	04/18/89			28.40		97.20		3.42		3.58	>	37.37
		62	01/09/90			93.20		320.00	>	3.43	>	6.47	>	37.10
	Charles Street Street	53	04/18/89			3.02		320.00		106.00				36.92
		57	07/11/89	0.554		89.00		309.00		3.47			>	36.88
		72	10/25/90				>	1000.00	>	0.97	>	10.67		

Table 16 (Cont'd)

	AVS	Ship-		Diff-				Language Control
Virus	No.	ment#	Date	rnt1.	IC 95	TC 95	AI 95	SI TAI
2000	2275	53	04/11/00	0.742	00 00 5	220 00 5	2 52	4 01 > 25 70
	2275		04/11/89		90.90 >		3.52	4.01 > 35.79
	3592	51	03/07/89		88.60 >		3.61	6.84 35.74
	6194	62	01/11/90		90.90 >		3.52	7.49 > 34.27
	6209	62	01/16/90		94.20 >		3.40 >	5.79 > 34.03
-	7087	72	10/25/90		9.03 >		11.07	6.67 > 33.38
	4796	61	12/06/89		30.90 >		10.35	10.03 32.83
	6208	62	01/16/90		94.00 >		3.40 >	5.94 > 32.42
	7910	75	11/06/90			3200.00 >	3.59	9.05 > 32.26
	6217 6211	62 62	01/16/90		94.40 >		3.39 >	5.69 > 31.74
	6223	62	01/16/90 01/16/90	0.022	92.60 > 298.00 >		3.46 1.08 >	6.40 > 30.64 2.07 > 30.15
	6205	62	01/11/90		222.00 >		1.44 >	5.92 > 30.06
	2506	64	03/13/90		9.43 >		33.94	6.68 29.71
	7083	72	10/25/90	0.919	85.70		3.68	6.21 > 29.69
	5998	61	02/01/90	0.633	3.02 >		10.61	9.75 29.64
	0206	4	12/05/89			1000.00 >	3.57	6.83 29.55
	6224	62	01/16/90		293.00 >		1.09 >	2.41 > 29.55
	5484	53	12/14/89		313.00	973.00	3.11	4.48 > 29.33
	5691	57	07/19/89		94.50 >		3.39 >	5.66 > 28.90
	2563	48	02/07/89		0.91 >		3.51	4.51 > 28.54
	4609	48	12/05/89		0.10 >		10.55	5.22 > 28.27
	0360	48	02/07/89		0.09	2.93	32.00	4.50 > 28.15
	5940	60	11/30/89		28.50	96.60	3.39	4.90 27.95
	6204	62	01/11/90		29.00	98.60	3.40	4.15 > 27.69
	6391	63	03/22/90			1000.00 >	3.50	4.95 27.54
	4425	44	03/29/89		29.50	103.00	3.49	2.20 > 27.43
	6207	62	01/11/90		279.00 >		1.15 >	4.09 > 26.05
	6580	64	03/15/90		93.90 >		3.41	4.62 > 25.97
	6220	62	12/05/90			1000.00 >	3.39	4.46 > 25.64
	2811	48	02/07/89		0.09 >		11.30	5.49 25.35
	7940	75	11/08/90	0.777	320.00	962.00	3.01	5.50 > 25.33
	6444	63	03/01/90		278.00 >		1.15 >	4.04 > 24.65
	7032	69	07/31/90		293.00 >		1.09 >	2.41 > 24.46
	4939	51	03/07/89		9.14	84.50	9.25	2.49 > 24.44
	4592	48	03/01/89		29.70 >		3.37	3.17 24.18
	7418	70	08/23/90		277.00 >		1.16 >	4.02 > 23.86
	5997	61	11/01/90		3.02 >		33.16	4.41 > 23.56
	5350	53	04/12/89		29.50	276.00	9.35	1.75 23.36
	6064	62	12/21/89		285.00 >		1.12 >	3.20 > 23.26
	2980	48	02/08/89		2.77 >		1.16 >	4.10 > 23.21
	4071	62	12/05/89			1000.00 >	3.13	4.35 > 22.39
	6200	62	01/11/90		302.00 >	320.00 >	1.06 >	1.79 > 22.36
	5495	53	04/18/89		2.71	189.00	69.80 >	3.71 > 22.03
	0361	48	02/07/89		0.03	0.57	19.20	3.93 > 21.60
	2906	53	04/11/89		94.50 >		3.39	2.51 21.14
	6226	62	01/16/90		291.00 >		1.10 >	2.55 > 20.41
	2812	48	02/07/89		0.01 >		1.11 >	2.87 > 19.34
	4822	48	11/28/89		31.10 >		10.28	3.49 18.49
	6988	68	09/13/90			1000.00 >	3.17	3.53 > 18.27
	8378	76	12/13/90		295.00	970.00	3.29	3.18 17.44
	6196	62	01/11/90		298.00 >		1.07	0.06 > 17.11
	4452	44	07/07/89		100.00 >		3.20	3.58 17.09
	6225	62	01/16/90		300.00 >		1.07 >	1.92 > 16.91
	6335	63	03/15/90			3200.00 >	1.08 >	2.22 > 16.87
	3612	32	11/03/88		293.00 >		1.09 >	2.45 > 15.95
	3450	32	04/19/89		29.70	95.80	3.22	1.71 15.85
	7023	69	09/06/90			1000.00 >	1.11 >	2.82 > 15.55
	2979	27	04/19/89		9.28	92.60	9.99	1.36 14.81
	7039	69	07/31/90		300.00 >		1.07 >	2.18 > 14.69
	7022	69	09/06/90		97.30	313.00	3.22	3.00 14.68
	4844	48	02/21/89		299.00 >		1.07 >	1.98 > 14.30
	4070	48	02/08/89		2.97 >		1.08 >	2.11 > 14.01
			, ,					

Table 16 (Cont'd)

Virus	AVS No.	Ship- ment#		Diff- rntl.	IC 95	TC 95	AI 95	sı	TAI
PT	7445	73	10/25/90	0.834	3010.00 >	3200.00 >	1.06 >	1.82	> 13.49
PT	6748	64	03/15/90	0.339	289.00 >	320.00 >	1.11	0.87	> 13.42
PT	6212	62	01/16/90	1.022	300.00 >	320.00 >	1.07 >	1.94	> 12.18
PT	6192	62	01/11/90	0.793	302.00 >	320.00 >	1.06 >	1.79	> 12.12
PT	5121	56	11/01/90		300.00 >	320.00 >	1.07 >	1.88	> 10.68
PT	4765	44	11/08/88	0.511	100.00	310.00	3.10	0.18	10.13
PT	6198	62	01/11/90	0.792	302.00 >	320.00 >	1.06 >	1.79 2	> 10.08
PT	4843	48	02/21/89	0.377	299.00 >	320.00 >	1.07 >	2.00	> 9.93

New Drugs with 50% Antiviral Reduction Levels: Out of the 6693 actual single drug tests, 364 new compounds demonstrated good antiviral activity, having antiviral reduction values equal to or better than 50%. This represents around 5.4% of the test compounds being active at this good antiviral reduction levels. These compounds are summarized in Table 17 according to the highest Total Antiviral Index (TAI). Four compounds (AVS-0303, 4851, 2320 and 4861) demonstrated the best TAI of >50% and SI is that ranged from 37 - 217. Sixteen (16) other compounds demonstrated moderate antiviral activity, having TAI's from 30 - 47% and SI's from 5 - 239. The rest (344 compounds) showed marginal antiviral activity with TAI's from 0.2 to 29% and SI's from > 1 to 95.

It is worthwhile to note (Table 17) that compounds received in shipment number 62 were mostly colored. Therefore those compounds appearing in the 50% active category from shipment number 62 should be interpreted with caution, since colored compounds create false positive readings with the MTT assay.

											50		
	AVS	Ship-	Test	Diff-									
Virus		ment#		rntl.	IC 50)	TC 5)	AI	50	SI		TAI
PT	0303	46	01/24/89	0.807	4.29	; >	320.00	>	75.	20 :	75.20	>	59.26
PT	4851	48	02/22/89		1.47	' >	320.00	>	217.	00 >	217.00		
	2320	53	04/11/89		2.50	5	172.00)	67.	30	36.50	>	52.20
PT	4861	48	02/22/89	0.401	6.80	>	320.00) >	47.	10 >	47.10	>	50.93
PT	4849	48	02/22/89	0.435	1.34	>	320.00) >	239.	00 >	239.00		
PT	4827	48	02/14/89	0.337	12.70	>	320.00	>	25.	10 >	25.10	>	43.84
PT	5242	52	03/08/89	0.798	14.50		320.00) >	22.				42.33
PT	5496	53	04/18/89	0.672	6.63		100.00		15.			>	40.80
	5451	53	04/18/89		2.5		76.90		30.				37.54
	4829	48	02/14/89		10.00		320.00			00 >			36.95
	6942	69	07/24/90		14.90		320.00			45 >			36.84
		60	01/23/90		0.63		10.00			34 >			36.45
	6945	69	07/26/90		16.50		320.00		19.				36.17
	1019	28	03/06/90		1.59		96.20		60.				35.94
	6943	69	07/24/90		60.60		320.00			28 3		>	32.49
	3181	29	05/17/89		1.65					40 70 >			31.66
	5498	53	04/18/89		30.00 19.20		320.00 269.00		14.		7.21	_	31.59
	1217 1214	52 52	03/15/89		20.50		211.00		10.			-	30.66
	5186	58	03/15/89 10/04/89		38.20		763.00		19.	_	10.31		29.76
	0111	9	12/07/89		22.50		830.00		36.			-	29.33
	4850	48	02/22/89		2.30		243.00		106.				29.33
	6603	64	04/18/90				1000.00			03 >	_		29.32
	5241	52	03/08/89		61.60					20 3			29.23
	7301	70	09/13/90				1000.00		16.				28.98
		61	11/29/89		6.3		320.00		50.		37.65		28.65
	2720	22	04/21/89		1.00		100.00		100.		67.70		28.46
	6745	64	03/08/90		51.30		210.00			10		>	28.40
	5869	60	01/23/90		61.10		758.00		12.		7.69		28.34
	7116	70	09/13/90				1000.00		12.		9.53	>	28.25
PT	4897	46	01/31/89		17.30		170.00		9.		5.27		27.87
PT	6714	67	07/12/90	0.660	42.30)	585.00)	13.	85	8.95		27.79
PT	5048	48	02/28/89		8.69	>	320.00	>	36.	80 3			27.34
PT	4763	44	12/12/89	0.778	32.00	>	1000.00	>	31.	25	0.23	>	27.27
PT	6412	66	07/06/90	0.561	6.0		74.70)	12.	43	8.81	>	26.81
	5855	60	01/23/90	0.688	74.80)	813.00)	10.	87	7.57		26.73
PT	6186	62	01/11/90	0.661	10.00		221.00		22.				26.26
PT	8377	76	12/13/90		37.30		255.00		6.			>	26.15
	7303	70	09/13/90		159.00		985.0		6.		3.87		25.87
	5075	48	03/01/89		48.10					65 :			25.83
	1992	56	06/14/89		57.70		320.00		5.				25.78
	8234	75	11/20/90		1.14		100.00		87.				25.09
	3180	29	05/17/89		0.7		100.00		141.				24.92
	8702	76	01/22/91		10.00		320.00			00 3			24.65
PT	2907	26	09/07/89	0.546	20.20)	210.00)	10.	38	6.50	>	24.44

Table 17 (Cont'd)

				-1					
Virus	AVS No.	Ship-ment#	Test Date	Diff- rntl.	IC 50	TC 50	AI	50	SI TAI
	6501	66	05/17/90		16.20	66.00		. 08	3.03 > 24.26
	4978	27	12/05/89		144.00	864.00		.02	3.65 > 24.04
	4943	51	03/07/89		1.79	22.20		40	6.49 > 24.01
	4785	46	01/12/89		1.00	29.10		.10 >	7.19 > 23.97
	8240	75 63	11/20/90		182.00	1990.00		.91	6.40 > 23.54
	6174 4106	62	01/09/90		136.00 >			.35 >	2.35 > 23.53
	4942	37 51	02/24/89		0.07 > 68.40	252.00		.25 > .68	4.25 > 23.24 1.93 > 23.05
	3964		03/07/89		2.68	24.50		. 16	4.58 > 22.98
	7439	33 73	03/22/89		153.00	638.00		.16	2.96 > 22.77
	3819	65	10/25/90 07/17/90		27.00	188.00		.97	4.06 > 22.75
	6473	66	07/10/90		8.61	171.00		91	11.31 > 22.68
	4934	51	03/07/89		2.43	20.50		. 43	4.48 > 22.59
	5060	48	02/28/89		262.00 >			.22 >	1.22 > 22.52
	0124	64	03/08/90		57.80			.53	3.84 22.45
	7349	70	08/16/90			1000.00		. 60	3.93 > 22.45
	5794	59	11/29/89		5.27	130.00		.74	5.15 > 22.31
	6177	62	01/09/90		247.00 >			.29	1.15 > 22.19
	8228	75	11/20/90		56.60	426.00	7.	.52	4.14 > 22.15
	6477	66	06/06/90		8.92	62.50		.01	4.56 > 22.06
PT	0148	2/67	10/04/90		0.29	2.90	9.	.99	4.88 > 22.01
PT	3621	32 .	12/20/89	0.673	63.40	528.00	8	. 33	4.66 21.94
PT	5201	58	07/26/89		78.10 >	320.00		.10 >	4.10 > 21.90
PT	8218	75	11/20/90	1.037	183.00	841.00	4	. 59	3.14 21.70
PT	5033	48	02/28/89	0.539	55.60	150.00	2	.70	1.10 21.68
	5230	52	03/15/89		72.70	259.00		. 56	1.23 21.64
	6422	66	07/06/90		49.10	241.00		.91	3.48 21.64
	4739	64	03/08/90		7.89	61.10		. 75	4.95 > 21.56
	8355	76	12/20/90		16.90	149.00		. 82	3.64 > 21.43
	5262	52	03/21/89		7.38	226.00		. 60	14.90 21.37
	7085	72	10/25/90		12.40	64.60		. 23	3.34 > 21.13
	6828	68	08/02/90		129.00	767.00		. 95	4.22 20.67
	7308	70	09/13/90			> 1000.00		.95	4.20 20.67
	3169	29	05/16/89		8.13	98.70		.10	8.04 20.60
	4875	46	12/20/89			1000.00		. 88	9.51 > 20.53
	5251	52	03/14/89		24.60 3.05	55.00 149.00		. 24	1.32 > 20.48 20.60 20.36
	4985 4809	51 46	03/14/89 01/17/89		42.50	301.00		. 07	3.53 > 20.27
	3064	28	04/21/89		5.03	22.90		. 56	3.28 20.24
	4459	45	11/23/88		76.00 >		× 4	.21 >	4.21 > 20.21
	8511	76	12/19/90			3200.00		.86 >	4.86 > 20.21
	8701	76	01/22/91		39.40			.13 >	8.13 > 20.17
	6497	66	05/17/90		4.83	40.40		. 35	4.72 > 20.04
	8237	75	11/20/90			3200.00		. 54	4.39 > 20.03
	7003	69	07/26/90		144.00			.22 >	2.22 > 19.93
	8259	76	12/11/90			3200.00		.94	4.68 19.80
PT	8233	75	11/20/90		367.00 >	3200.00	> 8	. 72	0.75 > 19.73
	8499	75	11/29/90		74.70 >			. 29	3.06 > 19.70
	4107	37	02/24/89		0.30 >			.08 >	1.08 > 19.43
PT	5450	53	04/12/89		4.47	27.40	6	. 14	3.57 > 19.39
PT	6977	68	08/02/90		28.90	210.00	7	. 27	5.37 19.33
PT	1654	46	01/24/89	0.803	6.23 >	320.00	> 51	. 30	1.06 > 19.28
PT	5069	48	11/28/89		41.20	204.00	4	.94	3.41 > 19.26
PT	8352	75	11/29/90		1.31	8.03	6	. 13	4.25 > 19.06
	8229	75	11/20/90	1.047	236.00	1450.00		. 16	3.05 > 18.87
	8281	75	11/28/90		54.10	191.00		. 53	1.80 > 18.48
	7449	71	07/31/90		71.30	184.00		. 58	1.63 > 18.38
	6102	62	12/28/89		106.00 >			.02 >	3.02 > 18.35
	6107	62	12/28/89		19.10	148.00		.76	3.28 18.32
	4241	46	01/24/89		3.99	23.20		.81	3.51 > 18.23
	5283	52	01/09/90		8.22	210.00		. 56	3.13 18.23
	6210	62	01/16/90		85.00			.77 >	3.77 > 18.16
	4715	43	12/13/88		212.00 >			.51 >	1.51 > 18.14
PT	5131	56	06/27/89	0.790	2.21	21.10	9	. 57	7.00 > 18.02

Table 17 (Cont'd)

	AVS	Ship-	Test	Diff-										
Virus		ment#	Date	rnt1.		IC 50		TC 50		AI 50)	SI		TAI
												-		
	6837	68	08/02/90			27.30		198.00		7.23	3	4.94		17.74
	7057	72	10/25/90			72.90	>	320.00		4.39		3.54		
	5483	66	07/17/90		<	0.03		0.27	>	8.33		5.73		
	7086	72	10/25/90			6.06		28.30		4.67		3.16		
_	6445	63	03/01/90			22.60		66.00		2.92		2.17		
	5643	57	08/16/89			52.70		280.00		5.32		3.61		16.92
	3678	32	11/10/88			1.46		10.00		6.86		3.92		
_	6978 6417	68 66	06/21/90			172.00 9.75	>	320.00 882.00	>	1.86		1.86		
	3168	29	07/06/90 05/16/89			3.02		100.00		90.49		5.95 3 33.10		
	6362	63	02/22/90			395.00		320.00		0.81		0.81		
	0919	52	03/15/89			65.70		186.00		2.83		1.59		
	3588	32	04/19/89			15.90		63.50		3.99		2.44		
	8376	76	12/13/90		•	17.30		110.00		6.37		3.84		
-	5501	53	04/18/89			3.08		65.30		21.20		15.60		
PT	2318	67	07/12/90			0.69	>	10.00	>	14.52		4.21		16.14
PT	5072	48	03/01/89			22.60		159.00		7.01		3.23	> :	16.07
PT	2604	19	12/21/88			25.50	>	100.00	>	3.91	\ >	3.91		
PT	6979	68	09/13/90	0.994		199.00		660.00		3.31	L	2.46		15.96
	6191	62	01/11/90			196.00		320.00		1.63		1.49		
	4222	48	02/08/89			155.00	>	320.00	>	2.07		2.07		
	4995	51	03/14/89	0.838		6.03		18.60		3.09		1.12		15.77
7.5	6873	69	09/05/90			81.30		213.00		2.61		1.85		
	8271	76	12/12/90			198.00		694.00		3.50		2.56		
	4860	48	02/21/89			285.00	>		>	1.12		1.12		
	7434 6442	70 63	10/02/90			198.00		642.00		3.24		2.34		
	6482	66	03/01/90 07/10/90			100.00		239.00		2.39		1.70 2.18		15.38 15.38
	2503	21	11/28/89			0.64		10.00		15.73		4.46		15.35
	4527	47		1.121		5.01		24.70		4.92		2.31		
	6903	69	07/19/90			222.00	>	320.00	>	1.44		1.44		
	5035	48	02/28/89			6.66		21.50		3.23		1.35		14.70
-	1980	27	05/16/89			2.59	>	100.00	>	38.50		28.30		14.44
	6031	61	11/09/89			82.30		262.00		3.18		2.20		14.42
PT	7414	70	08/23/90			202.00		986.00		4.88		3.16		14.40
PT	4590	61	12/06/89			0.53		2.62		4.91	L	2.46		14.26
	8224	75	11/20/90			189.00		779.00		4.13		1.70		
	7055	72	10/25/90			63.00		210.00		3.33		2.46		
71.5	7072	72	10/25/90			206.00		646.00		3.13		2.27		
	6249	1P	11/14/89			72.50		269.00		3.71		2.55		14.05
_	7399	70	09/27/90			188.00		804.00		4.27		2.74		13.90
	7042	69	07/31/90			70.60		208.00		2.94		2.15		13.81
	5503	53	02/13/90			100.00		210.00		2.10		1.55		13.80
	5207 4853	58 48	11/30/89 02/22/89			9.12 1.73		20.30 7.72		2.23		1.58 : 2.52 :		
	6029	61	02/22/89			60.60		210.00		3.47		2.56		13.65
	7094	72	10/25/90			63.80		259.00		4.21		2.67		
	6160	62	01/04/90			5.83		18.10		3.10		1.91		
	2453	18	11/28/89			5.90		20.30		3.44		2.45		13.44
	1774	56	06/14/89			300.00	>	320.00	>	1.07		1.07		
	7033	69	07/31/90			61.00	-	241.00		3.95		2.79		13.42
	5905	61	11/29/89			5.66		19.20		3.40		2.27		13.22
PT	4795	46	01/12/89	0.885		69.00		275.00		3.99	•	2.16		13.19
	8312	76	12/12/90			718.00		1870.00		2.61		1.69		13.14
	7484	73	10/30/90			181.00		711.00		3.92		2.84		13.00
	8263	76	12/11/90			47.90		164.00		3.42		1.63		13.00
	4770	44	11/08/88			60.10		202.00		3.39		2.37		12.94
	6970	68	08/02/90			27.70		167.00		6.04		3.35		12.85
The state of the s	4825	48	02/14/89			68.40		240.00		3.51		1.57		12.84
Laborator .	8358	76	12/20/90	0.937		54.90		204.00		3.71		2.53		12.83
	4919	61	12/06/89			53.10		189.00		3.56		2.33		
	1355 4278	64	05/03/90			231.00 9.75		1000.00		4.33		2.85		
	4769	44	12/06/89 11/08/88			58.20		51.60 192.00		5.30		2.51		12.66
F.4	4,03		11,00,00	0.341		30.20		192.00		3.30	•	2.20	•	12.63

	3110	ah !		2166										
Virus	AVS	Ship-	Test	Diff-	TO 50					AT 50				
ATLUB	NO.	ment#	Date	rnt1.	IC 50		TC !	50		AI 50		SI		TAI
PT	5191	58	11/30/89	0.920	0.93		6.	22		6.78	1	A 7A		12.54
	2290	71	07/31/90		496.00	>	320.		>	0.65				12.52
	7430	70	09/27/90		289.00		660.			2.28		1.69	>	12.44
	2450	18	11/28/89		20.80		100.		>	4.81		2.56		12.40
PT	4281	61	12/06/89		2.35		12.	00		5.11		2.93		12.39
	6904	69	07/19/90		24.30		66.	00		2.72				12.37
	6601	64	04/18/90		204.00		897.			4.40			>	12.31
	4891	46	12/20/89		0.24		0.			2.87		2.11		12.16
	2151 6436	52 66	03/15/89 07/06/90	0.881	80.90		314.			3.88 2.63		1.82		12.14
	8219	75	11/20/90		24.20 163.00		63.			2.22		1.89		12.11
	5837	59	11/30/89		24.40		61.			2.52				12.03
	6236	GABSN			3080.00	> 3			>	1.04				12.02
	1736	45	11/16/88		82.10		320.			3.90				12.01
PT	6379	63	02/27/90		217.00	>	320.	00	>	1.47		1.47	>	11.96
	5482	66	05/08/90		21.00		86.			4.12			>	11.78
	7935	75	11/08/90	0.927	157.00		609.			3.88		2.53		11.71
_	6234	GABSN	02/08/90		1600.00					2.00				11.69
	6041 6753	61	11/09/89		282.00					1.13			>	11.64
	5500	67 53	07/12/90 04/18/89		255.00 : 1.90	> 1	6.4		_	3.38		3.40		11.59
_	8260	76	12/11/90		239.00	> 1			>	4.18				11.53
	0053	64	03/08/90		100.00		265.			2.65				11.49
	7373	70	09/18/90		201.00		614.			3.05				11.39
PT	6337	63	03/20/90	0.945	19.70		67.			3.42	2	2.52		11.38
	4753	44	12/14/89		2.92		17.			6.01		3.53		11.34
	4992	61	12/06/89		200.00		638.			3.19		2.28		11.34
	4757	44	12/14/89		299.00		660.			2.21			>	11.33
	6071 3112	62 28	12/21/89		6.58 8.70		21.			3.26		2.39		11.30
	7488	73	04/25/89		370.00	> 1			>	2.70				11.23
	6182	62	01/09/90		160.00		320.			2.00				11.11
	8272	76	12/12/90		766.00					4.18				11.10
	6213	62	01/16/90		21.40	_	62.			2.92		1.96		11.00
PT	5136	57	07/11/89		81.90		210.	00		2.56		1.89		10.99
	5452	53	04/18/89		5.47		18.			3.34				10.97
	7415	70	08/23/90		170.00		320.		>	1.88			>	10.97
	6842 7942	68 75	08/22/90	0.769	56.60 203.00		143.0 482.0			2.53		1.36		10.93
	3491	65	11/08/90 04/17/90		190.00		320.		`	1.69				10.82
	8250	76	12/11/90		75.60		359.			4.75		2.67		10.68
	6075	62	12/21/89		68.40		207.			3.02			>	10.65
	6201	62	12/05/90		192.00		320.		>	1.66		1.66	>	10.57
	6443	63	03/01/90	0.583	77.10	>	320.			4.15	5	2.72	>	10.57
	2573	57	06/27/89		286.00		320.			1.12				10.54
	6583	64	03/15/90		320.00		320.0		>	1.00			>	10.54
	6011	61	02/01/90		275.00		623.			2.26		1.43		10.47
	0002 5189	46 58	01/24/89 11/30/89		187.00		320.	20		1.71				10.46
	4768	44	11/08/88		66.80		187.			2.80		1.78		10.27
	8374	76	12/13/90	0.955	566.00		890.			3.34		2.18		10.26
	3613	32	11/08/88		61.40		320.		>	5.21			>	10.23
PT	4105	48	02/08/89		7.37		22.			2.99)	1.51		10.07
	1159	52	08/23/89	0.697	60.40		204.			3.38	}	2.18		10.02
	5917	60	01/23/90	0.734	2.68		7.1			2.92		2.00		9.97
	5520	56	06/14/89		236.00		320.			1.35		1.35		9.91
	4075	65	05/15/90		97.80		320.		>	3.27		2.56	>	9.82
	7469 5128	73 56	10/30/90		185.00 10.00		603.0			3.25		2.06		9.82
	1733	45	06/27/89 11/22/88		69.80		21.0			3.01		1.55	-	9.53
	7930	75	11/08/90		691.00		410.			3.49		2.24		9.51
	6127	62	12/28/89		19.40	_	58.			3.01		1.84		9.48
	7944	75	11/08/90		534.00	2	000.			3.75		0.17		9.48
PT	4591	63	02/13/90	0.615	0.01		0.0			3.33		2.30	>	9.47

Table 17 (Cont'd)

	AVS	Ship-	Test	Diff-										
Virus		ment#		rntl.		IC 5	0	TC 5	0	AI 50		SI		TAI
рπ	5538	56	08/09/89	0.658		256.0	0	663.0	^	2.59		1.92		9.41
	6584	64	04/18/90					1000.0		1.59	>	1.59	>	9.27
	6449	63	03/01/90			264.0		320.0		1.21		1.21		9.14
PT	5448	53	04/12/89			5.8		19.8		3.41		1.34		9.08
PT	1337	67	05/30/90			79.0	0	234.0	0	2.96		2.11		9.02
	5988	61	12/01/89			7.4		22.8		3.05		2.12		8.96
	5765	59	01/23/90	0.765		257.0	_	320.0		1.25	>	1.25	>	8.86
	8328	76	12/13/90			258.0		642.0		2.49		1.80		8.67
	5539	56	08/09/89			2.3		7.3		3.18	_	2.14		8.62
	7049 7040	69	09/11/90			71.4		1000.0 305.0		1.66	>	1.66	>	8.53
	7945	69 75	07/31/90 12/04/90	1 204		854.0		1570.0		4.27		2.48	•	8.47
	4800	46	01/17/89			245.0		320.0		1.31	>	1.31		8.45
	6181	62	01/09/90			320.0		320.0		1.00		1.00		8.43
	3374	31	12/20/88	0.993		84.4		100.0		1.18		1.18		8.37
PT	6839	68	08/22/90	0.849		75.9		225.0		2.97		2.14		8.37
PT	1750	45	11/23/88	0.722		202.0		320.0	0 >	1.58	>	1.58	>	8.26
	8251	76	12/11/90			91.8		296.0		3.23		2.16		8.21
	4383	43	12/14/88			194.0		320.0		1.65		1.65		8.17
	3550	31	03/29/89			94.7		100.0	_	1.06		1.06		7.99
	6521	66	07/10/90			250.0		320.0		1.28	>	1.28		7.95
	8230 8270	75 76	11/20/90			735.0		1000.0		1.41 2.55		0.68		7.90 7.85
	2543	76 21	12/12/90 11/28/89			68.4		100.0		1.46	>	1.46		7.77
	6722	67	06/05/90			203.0		320.0		1.57		1.57		7.77
	6440	63	03/01/90		<	1.0		9.5		9.51		1.88		7.67
	7074	72	10/25/90			320.0		660.0		2.06		1.26		7.64
PT	3560	31	03/29/89			279.0		320.0		1.15		0.64		7.60
PT	6730	67	06/05/90	0.636		173.0		320.0		1.84		0.39	>	7.56
PT	3425	32	04/19/89	0.515		30.5		46.9		1.54		0.83		7.45
	2579	19	11/28/89			281.0		320.0		1.14	>	1.14	>	7.38
	7487	73	10/30/90			212.0		622.0		2.93		1.08		7.34
	2575	57	12/05/90			235.0		940.0		4.00		1.87		7.31
	5842 2590	60 65	11/30/89 04/12/90			97.8 227.0		251.0 821.0		2.56 3.62		1.79	>	7.19 7.15
	3565	32	04/05/89			7.7		19.4		2.49		1.65		7.06
	6983	68	06/21/90			187.0		320.0		1.71	>	1.71	>	7.06
	4277	42	11/28/89			7.9		10.0		1.27		1.27		7.04
PT	1850	67	05/31/90	0.715		189.0	0	497.0	0	2.63		0.40		7.03
PT	5835	59	11/29/89			299.0	0	300.0	0	1.00		0.67		7.02
	4424	44	03/29/89			30.0		60.6		2.02		1.07		6.96
	6062	62	12/21/89			205.0		320.0		1.56		1.56		6.86
	7424	70	08/23/90			185.0		320.0		1.73	>	1.73	>	6.82
	6976 7048	68 69	09/13/90 09/11/90			72.7		173.0 165.0		2.38		1.38		6.67
	4223	63	02/13/90			0.2		0.6		2.10		1.08		6.39
5.0.0 To	5525	56	06/14/89			270.0	_			1.18	>	1.18	>	6.38
	2580	19	11/28/89			8.1				1.23		1.23		6.35
	6607	64	04/18/90			9.0		18.5		2.05		1.29		6.28
	6138	62	12/28/89			20.6		59.5		2.89		0.36		6.23
	6374	63	03/20/90			259.0				1.23	>	1.23	>	6.15
	8698	76	01/22/91	0.858		24.6	0	83.8		3.41		1.76		6.12
	3107	28	04/25/89			28.0		38.2		1.37		0.77		6.11
	0347	62	01/18/90			2.5	_	6.4		2.49		1.54		6.04
	4611	65	05/15/90			133.0				2.41		0.12	>	5.88
	6861	68	08/02/90	0.640		320.0		710.0		2.22		1.57		5.88
	8373 6103	76 62	12/13/90 12/28/89			9/8.0		3200.0		3.27 2.19		1.02	-	5.76 5.71
	6055	62	12/21/89			1.0		3.7		3.71	>	1.92	>	5.60
	8309	76	12/12/90					3200.0		1.66		1.66		5.48
	6222	62	01/16/90			239.0				1.34		1.14		5.45
PT	6221	62	01/16/90	1.037		255.0				1.26		0.50		5.38
PT	7405	70	08/23/90			247.0				1.29	>	1.29		5.36
PT	5823	59	11/29/89	0.785		94.6	0	186.0	0	1.96		0.96		5.35

	AVS	Ship-		Diff-		10.00		Part No.					17.07		
Virus	No.	ment#	Date	rntl.		IC 50		TC 50		AI	50		SI		TAI
DO	7375	70	09/21/00	0 072		90.90		320.00		•	52		0.14		5.19
	8332	76	08/21/90 12/13/90			245.00		531.00			17		1.21		5.18
	7385	70	08/21/90			82.20		184.00			24		1.42		5.16
	6189	62	01/11/90			224.00					43		0.45	>	5.09
	5489	66	05/08/90			263.00					22	>	1.22		5.04
	6843	68	08/22/90			72.00		175.00			43		1.13		4.98
	7408	70	08/23/90			305.00	>		>		05	>	1.05	>	4.96
PT	3561	31	03/29/89			300.00	>	320.00	>	1.	07	>	1.07	>	4.87
PT	6180	62	01/09/90	0.588		238.00	>		>		34	>	1.34	>	4.84
	6308	63	02/15/90			9.30		19.50			10		1.42		4.81
	3098	28	04/25/89			28.30		42.60			50		0.82		4.79
	5818	59	11/29/89			74.90		110.00			48		0.69	>	4.76
	7047	69	08/02/90			85.50		194.00			27		1.53		4.58
	5506	53	04/18/89			31.30		59.50			90		1.26		4.42
	8311 2363	76 64	12/12/90			819.00 29.80		1830.00 78.80			23		0.01		4.39
	5459	54	03/13/90 05/05/89			9.57		24.90			60		1.16		4.15
	6334	63	02/20/90			77.60		182.00			34		0.41		4.07
	7485	73	10/30/90			235.00		723.00			08		0.87		3.96
	7004	69	09/06/90			308.00		550.00			78		0.90		3.89
	3110	28	04/25/89			25.20		46.30			84		0.99	>	3.87
	7406	70	09/27/90			320.00		443.00			38		0.75		3.63
	4074	48	12/07/89		<	1.00		2.71			71	>	1.69		3.55
PT	5003	45	03/01/89			305.00	>	320.00	>	1.	05	>	1.05	>	3.53
	7071	72	10/25/90	1.089		246.00		495.00		2.	01		0.94	>	3.39
	7320	70	09/13/90			320.00		456.00			43		0.76		3.35
	6792	67	06/12/90			3.10		5.77		1.	86		1.12		3.31
	8541	76	01/03/91	0.808		28.30		59.50			10		0.03		3.27
	8349	75	11/29/90	0.890		293.00	>		>		09		0.64	>	3.13
	3589	32	04/19/89			26.10		30.80			18		0.78	_	3.04
	8330	76	12/13/90				>	3200.00	>		17		1.09	>	3.04
	8236 7417	75 70	11/20/90 08/23/90			94.00 254.00		320.00 320.00			40		0.62		2.95
	4864	65	05/15/90			267.00					20		0.34		2.93
	6243	1P	11/14/89					1000.00			19		1.14		2.85
	7925	75	11/08/90					3200.00			32		0.79		2.83
	8366	76	12/13/90			260.00		477.00			84		0.76		2.73
	8685	76	01/22/91			100.00		181.00			81		1.11		2.73
PT	7383	70	08/21/90			245.00	>	320.00	>		31		1.07	>	2.68
PT	7391	70	08/21/90	0.994		80.00		185.00		2.	31		0.40		2.54
	5005	45	03/01/89	1.078		320.00					00		1.00	>	2.48
	2572	65	04/12/90			306.00					05		1.05		2.47
	6923	69	07/24/90			299.00	>		>		07	>	1.07	>	2.39
	8368	76	12/13/90			300.00		720.00			40		0.82		2.39
	7350	70	08/16/90	0.935			>	1000.00	>		80		0.94	>	2.28
	8375	76	12/13/90			301.00		531.00			77		0.74		2.28
	7379	70	09/18/90					1000.00			09		1.09		2.25
	7410	70	08/23/90					320.00			00	>	1.00		2.16
	8318 6560	76 66	12/12/90 07/10/90					3200.00 1000.00			35 13		0.33		2.02
	5983	61	02/01/90			9.38		16.30			74		0.88		1.97
	8262	76	12/11/90				>	1000.00	>		17		0.90	>	1.82
	6153	62	01/04/90			1.00		2.10			10		1.55		1.68
	8372	76	12/13/90				>	3200.00	>		20		0.44		1.68
	7368	70	08/16/90					1000.00			09		0.74	>	1.23
	7017	69	09/06/90			289.00		460.00			59		0.54		1.22
	7011	69	09/06/90				>	1000.00	>		12		0.63		1.21
PT	7386	70	08/21/90	0.853		938.00	>	1000.00	>	1.	07		0.78		1.20
	7403	70	08/23/90				>	1000.00	>		02		0.69		0.85
	3101	28	04/25/89			32.00		39.40			23		0.66		0.66
	5405	53	03/28/89		<			1.73	>		73	~	1.00		0.62
	6215	62	01/16/90			239.00		100.00			42		0.12	>	0.25
PT	3105	28	04/25/89	0.861		32.00		35.90		1.	12		0.67		0.24

New Drugs with 25% Antiviral Reduction Levels: Of the 6693 actual single drug tests, 560 new compounds demonstrated moderate antiviral activity, having antiviral reduction values equal to or better than 25%. This represents around 8.4% of the test compounds being active at this marginal antiviral reduction level. In general, when compared to the 95% and 50% antiviral activity categories, the compounds in this (25%) category do not appear to have any significant antiviral promise and probably do not need to be presently confirmed any further.

4.1.6.3. Confirmatory Assays:

Some of the compounds were sent to us in more than one separate drug shipment. These compounds were tested more than once. Data from the confirmatory assays are summarized in Table 18. If a compound showed >50% reduction in CPE during this contract period, then it was considered a candidate for confirmatory testing. The confirmatory tests are from active compounds picked up by both the VR and MTT assay testing. Out of 326 confirmatory tests, 214 compounds were confirmed active during this reporting period and the remaining 112 compounds gave conflicting results. The criteria for activity is that the confirmatory test has to show $\geq 25\%$ reduction in CPE. Failure to confirm the activity in these compounds was probably due to differences during the assay conditions:

- 1) In confirmatory assays the concentration range is adjusted to a more accurate semilog scale to maximize the TAI window.
- 2) Differences in the "differential" of the two runs can cause the compound to read positive or negative, falsely. The variability in the differential can cause false positive or negative bias to be introduced into the calculations, thus reflecting the variability in the maximum activity of the compound.
- 3) The metabolic rate, cell and virus load/well, age, and passage number of the cells may cause the above observed variability in the confirmatory results.
- 4) Problems associated with stability and storage of the compound (i.e., different lot numbers, solubility, light sensitivity, hygroscopic, etc.).
- 5) Problems associated with technical execution of large numbers of plates by different technicians.

During this reporting period the overall confirmatory rate against PT was 66%. The conflicting results should be retested at a later date based on the availability of the compound.

4.1.6.4 Recommendations of PT-Actives Based Upon the In Vitro Results with MTT Assay (Vero Cells).

Based upon the in vitro results with the MTT assay (Vero cells) we recommend the following:

- 1) Compounds with the highest TAI in the 95% activity category that have confirmed results with the exception of "colored" compounds should receive the highest priority for further profile studies and in vivo animal testing.
- 2) Compounds with the highest TAI in the 50% activity category that have confirmed results with the exception of "colored" compounds should receive the next highest priority for further profile studies and *in vivo* animal testing.

Table 18

Confirmatory Assays for Compounds Active Against Punta Toro (PT)

< ∪ ⊢		* *	+	+	+	•	+	•	+	+	٠	+	•	•	•	+	•	•	•	+	•	+	•	٠	•	•	•	•	•	+	•	•
Assay Type		ĒĒ	36	w	S	3	TIM	H	Ħ	*	H	35	9	SPE	3	SPE CPE	H	Ħ	Ħ	SPE	2	9 8	SPE	2	CPE	SPE CPE	Ħ		Ħ	SPE CPE	CPE	Ħ
IAI		10.46	4.30	0.00	1.20	0.30	11.49	0.0	2.43	2.00	4.20	2.40	0.30	0.00	0.03	1.90	1.12	0.31	0.0	3.00	0.70	0.00	0.05	0.00	0.00	0.00	1.15	0.0	5.58	.9	0.00	4.20
5		1.71 > 0.00 >	17.60	2.60	3.20	0.00	1.70	9.0	0.00	2.00	0.00	13.30	0.00	0.00	0.00	4.80	0.00	0.0	0.00	32.00	0.00	1.2	0.00	0.0	0.00	0.0	0.00	0.00	0.0	6.20	0.0	0.00
A1 95		0.00	3.20	3.20	1.0	0.0	0.00	0.0	0.0	0.10	0.00	3.20	0.00	0.0	0.00	1.00	0.00	0.0	0.00	0.0	0.0	1.00	0.00	0.00	0.0	0.00	0.0	0.0	0.0	3.20	0.00	0.00
25 25		320.00 1000.00	320.00	320.00	100.00	32.00	320.00	320.00	320.00	10.00	100.00	32.00	10.00	10.00	10.00	320.00	320.00	100.00	864.00	320.00	320.00	320.00	320.00	320.00	320.00	100.00	100.00	1000.00	3200.00		320.00	
10 %		0.00	100.00	100.00L	100.001	0.00	0.00	0.00	0.00	100.00	0.00	10.00	0.00	0.00	0.00	320.00 >	0.00	0.00	0.00	0.00	0.00	320.00 >	0.00	0.00	0.00	0.00	0.00	0.00	0.00	100.00	0.00	0.00
AI 50		1.71	17.60 -	2.8	3.10 ~	0.00	5.65	9.0	0.00	- 09.0	0.0	13.30 ~	0.0	0.0	0.00	4.82 >	0.0	0.0	0.00	32.00	0.0	5.46	0.0	0.0	0.0	0.0	0.0	0.00	0.0	20.00	0.0	0.00
10 50		320.00 >	320.00	100.00	100.00	32.00	265.00	105.00	71.20	10.00	7.50	32.00	10.00	10.00	10.00	320.00	99.99	100.00	175.00	320.00	100.00	320.00	100.00	320.00	32.00	32.00	100.00	1000.00	1600.00	320.00	320.00	237.00
10 50		187.00 >	18.20 >	37.94	32.00	0.00	100.00	0.00	0.00	16.00 ~	0.00	2.40	0.00	0.00	0.00	07.99	0.00	0.00	0.00	10.00 >	0.00	130.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	16.00 ~	0.00	0.00
A1 25		2.46	1000.00	4.10	4.10	0.00	3.47	0.0	2.01	3.20	0.00	32.00	0.0	0.00	0.00	0.31	0.0	0.00	0.00	320.00	0.00	2.67	0.00	0.00	0.00	0.00	. %	0.0	4.20	10.00	0.00	0.00
10 25		320.00 >	320.00	99.99	99.00	32.00	179.00	28.70	45.40	32.00	3.50	32.00	10.00	10.00	10.00	100.00	34.50	93.20	67.60	320.00	99.00	160.00	90.99	160.00	16.00	16.00	100.00 >	517.00	90.099	100.00	10.00	111.00
10.25		130.00 >	0.32	16.00	16.00	0.00	51.60	0.00	21.10	10.00	0.00	- 1.00	0.00	0.00	0.00	320.00	0.00	0.00	0.00	1.00 >	0.00	90.09	0.00	0.00	0.0	0.0	91.90	0.00	157.00	10.00	0.00	0.00
Diff.		0.760	*	≤	≨	S	0.977	0.630	1.500	K	0.818	¥	×	K	¥	×	0.833	0.695	90.00	4	¥.	Y.	¥	¥	¥	¥	0.850	0.646	1.044	¥	¥	0.788
14		104		:		:	NEW CEN	8	162	:	PDA	:	:	:	:	:	95	8	E	:	:	:	:		:	:			2XR	:	:	99
Test		01/24/89 OE0 12/20/89 TOA	09/19/86	05/12/87	10/17/86	05/20/88	03/08/90	05/01/90 VSD	10/04/90 291	10/23/86	03/28/89 PDA	09/19/86	11/18/86	08/21/87	09/21/87	09/19/86	05/23/89	08/23/89 RBD	05/30/90	98/11/60	11/18/86	05/12/87	08/21/87	09/18/87	05/04/88	05/20/88	11/30/89 SNL	04/10/90	11/01/90	10/23/86	05/20/88	01/24/89 OEP
Ship- ment	PT	33	-	_	2	~	ઢ	ઢ	z	2	53	-	_	_	_	-	_	_	29	_	-	_	_	_	_	-	_	9	_	2	7	94
AVS S	** VIRUS P	0002	0015	200	0033	0033	0053	0053	0053	9000	9000	9900	8900	8900	8900	7800	7800	7800	7800	7600	7600	7600	7600	7600	7600	7600	7600	7600	7600	9600	9000	900
						4																										

Table 18 (Cont'd)

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Assay Type	Ė	H	HTT	HT	CPE	CPE	Jo.	2 6	1 1 1 1 1 1 1 1 1 1	H		H	TIM	E	SPE SPE	CPE	CPE	į	ב ב ב	e Co			ě	, w		8	8	TH	SPE CPE	B	35	3	TH	Ħ		H	CPE	S S	H.	
TAI	1.02	5.3	10.24	29.33	2.00	0.00	200	5 6	3	8:8	0.00	22.45	1.44	3.96	1.50	0.55	0.00	700	0.00	0.40	18.23	3.89 22.01	1.70	1.50		2.10	0.35	0.05	1.90	0.00	2.70	1.40	19.78	25.19	28.55	8.69	2.70	2.35	0.27	
ïs	0.00	0.0	0.0	11.42 >	10.00	0.00	5	8.6	9.0	0.0	0.00	3.84	0.00	0.00	4.38	0.33	0.00	00 000	00.00	3.20	4.61 ×	0.0 4.88.4	07.7	2.40		07.9	0.38	0.0	2.30	0.0	7.40	2.3	3.79 >	9.60	6.83	2.63	1.50	1.40	0.00	
A 98	0.00	0.0	0.00	0.00	0.00	0.00	8	3 8	0.00	0.00	0.00	0.00	0.00	0.00	0.00	1.25	0.00	200				88		0.0		9.	0.0	0.00	9.1	0.0	1.00 >	1.00	0.00	0.00	3.57	0.0	10.00	31.30	0.00	
7C 9S		100.00			32.00	100.00	100 00	9.00	30.00	96.60	8.00	320.00	1000,00	1000.00	320.00	275.00	275.00	730	350.00	2.00	3.20	10.00	M			100.00	100.00	100.00	320.00	320.00	320.00	320.00	320.00	1000.00	1000.00	1000.00	320.00	100.00	10.00	
36 DI	0.00	0.00	0.00	0.00	0.00	00.00	000	9.0	0.0	0.00	0.00	0.00	0.00	0.00	0.00	220.00 >	0.00		9.	8:	0.00	° ° °	^ 00 O	0.00	1	. 100.00	0.00	0.00	- 320.00 >	0.00	- 320.00 >	320.00 >	0.00	0.00	280.00 >	0.00	. 32.00	3.20	0.00	
A1 50	0.00	0.0	0.00	36.85	10.00	0.00		8 6	0.00	0.00	0.00	5.53	0.00	0.00	4.38	0.50	0.00	757	00.00	3.20	98.4	8.8	07.7	2.40		6.41	0.38	0.0		0.0	7.40	5.60	3.79	11.25	10.29	6.24	1.45	1.43	0.00	
TC 50	265.00	100.00	563.00	830.00	32.00	66.00	44 00	3.5	34.00	8.9	59.50	320.00	83.00	278.00	320.00	86.00	86.00	9	20.00	00.	3.20	2.50 2.80 2.80	320.00	320.00		100.00	32.00	82.50	320.00	320.00	320.00	320.00	320.00	8	871.00	585.00	3.20	1.00	2.20	
10 50	0.00	0.0	0.00	22.50	3.20	0.00	0	8 8	0.0	0.00	0.00	57.80 >	00.0	0.00	73.00 \$	171.50	0.0		0.32	0.32	^ 99.0 	0.0 0.0 0.0	73.00 ×	134.60 >		15.60	85.00	0.00	141.00	0.00	43.00 >	57.00 >	84.40 >	70.30	9.60	93.80	2.20	o.3	0.00	
AI 25	0.00	0.0	7.46	19.76	32.00	0.00		8 6	0.00	0.00	0.00	5.56	0.0	0.00	22.90	0.44	0.0	767	213.00	1.50	6.89	8.94	26.70	4.90		9.70	0.54	0.00	21.00	0.00	10.00	10.00	6.9	13.00	13.68	4.50	2.30	1.29	0.00	
70 25	112.00	88.70	183.00	257.00	32.00	32.00	42 00	36.00	16.00	69.00	39.30	222.00	50.30	181.00	320.00	56.80	56.80	00	9.0	0.32	30.0	1.42	320.00	320.00		99.99	16.00	26.70	210.00	320.00	> 320.00	320.00	320.00 >	764.00	578.00	247.00	2.30	99.0	0.94	
10 25	0.00	0.0	24.50	13.00	1.00	0.00	000	9.0	0.00	0.00	0.00	00.07	0	0.00	14.00 >	130.00	0.00		0.36	0.21	0.44	0.22	12,00 2	66.00		6.80	96.00	0.00	10.00	0.00		~ 32.00	46.00 >	35.70	42.30	24.80	1.00	0.51	0.00	
Diff.	0.808			0.772	¥	×					0.687	97.0			×		¥	•	S			1.514	4 2					0.646	¥	KA	¥	X		0.782			¥		0.747	
¥	80 018				98	78	87				89 RBF		90 VSF		98	88	88					25 25 29 25 29 29 29 29 29 29 29 29 29 29 29 29 29 2	98	88		98		89 RBG	98	88	98	98				SO WEK	98	28	89 RBH	
Test	05/10/89	08/23/89	12/07/89	12/07/89	09/19/86	08/21/87	78/1/187	7/40	06/13/88	05/23/89	08/23/89	03/08/90	05/01/90	04/11/70	10/27/86	05/31/88	05/20/88	707 007 00	10/61	08/21/87	08/23/89	10/04/90	10/30/86	05/18/88		10/21/86	05/20/88	08/23/89	10/28/86	05/20/88	10/28/86	10/30/86	05/10/89	12/05/89	12/05/89	05/30/90	10/03/86	09/24/87	08/23/89	
Ship-	24	75	0	•	_	-				- - ,	-	3	3	8	4	4	4	c	y (7	~!	2/67	•	· 10		~	~	~	4	4	10	5	'n	4	4 !	29	_	-	_	
AVS	1110	1110	0111	1110	0113	0113	0112	2 .	515	0113	0113	0124	0124	0124	0136	0136	0136	0.00	2	0148	0148	0148	0197	0197		0500	0500	0500	0202	0205	9020	9020	0206	9020	9020	9020	0215	0215	0215	

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	Type	EE	CPE	_		H	HTT	HTT	TTH	HT	H	MTT	8	8	H	H	H	Ħ	TTH	Ħ	MIT	H	5	5	5	E		TH.	E E	E		Ħ	Ę		
	IAI	38.83	2.30	2.00	0.07	59.26	5.49	16.85	6.04	28.15	8.9	0.00	1.10	1.00	21.60	0.60	2.03	0.07	16.54	5.57	4.41	33.92	3.40	8.	1.00	31.88	3	31.42	3.5	4.01	0.00	0.00	8.6	7.05	
	28	14.00 >	3.20	2.00	0.00	75.20 >	0.00	0.00		4.50 >	0.00	0.00	3.1	2.40	3.93 >	0.00	0.00	0.0	1.59 >			15.12 >	3.60	0.20	1.40	8.8	3	7.21	2.24 >	6.00	8 8	0.0	8:	11.7	
	A1 95	0.00 > 2.61 >	10.00	3.10	0.0	0.00	0.0	0.0	8.	32.00	0.00	0.00	0.00	3.20	19.20	0.00	0.00	0.0	0.00	0.00	0.00	0.00	100.00	3.20	0.0	8.8	3	0.0	8.	8.8	88	0.00	8.8	8.	
	20 21	320.00	100.00	10.00	100.00	320.00	320.00	32.00	25.20	2.93	9.	1.00	0.32	0.32	0.57	0.29	0.29	0.32	332.00	309.00	1.00	100.00	320.00	32.00	320.00	320.00	350.00	320.00	320.00	32.00	320.00	1000.00	320.00	840.00	
	26 25	0.00 >	10.00	3.20 >	0.00	0.00	0.00	0.00	0.00	0.0	0.00	0.00	0.00	0.10 -	0.03	0.00	0.0	0.00	0.00	0.00		0.00	3.20	10.00 >	0.00	8.0	8.5	0.00		0.00	000	0.00	• 6 • 6 • 6	8.	
	AI 50	14.00	100.00	20.00 -	0.00	75.20	0.00	0.00	5.49	10.80	0.00	0.0	1.7	8.00	5.76	0.0	0.0	0.00	2.63	0.00	1.11	60.63	3.60 -	0.23 ~	4.60	7.20	3	14.00	5.24	8.0	800	0.0	8.8	8.%	
	TC 50	320.00 >	100.00	10.00	0.87	320.00 >	100.00	7.99	6.45	77.0	0.29	0.93	0.32	0.32	0.00	90.0	90.0	0.08	186.00	207.00	1.8	96.20	3.20	0.32	320.00	218.00	9. 9.	269.00	320.00 >	32.00	320.00	320.00	320.00	254.00	
	10 50	22.90 > 13.90 >	1.00	0.50 >	0.00	4.25 >	0.00	0.00	2.59	9.0	0.00	0.0	0.50 >	90.0	0.01	0.00	0.0	0.00	65.70		0.00	1.59	0.0	1.42	< 00.69	30.00			^	0.00					
	AI 25	30.10	10.00	3.10	0.00	320.00	1.28	24.20	3.2	9.21	3.44	0.70	3.20	3.10	6.82	0.0	2.47	0.00	34.30	1.50	8:	23.19	3.61	0.25	3.10	58.00	9.0	11.20	12.36	1.63	0.0	0.0	0.0	3.32	
	25 21	320.00 >	3.20	1.00	09.0	320.00 >		5.59	3.8	0.18	0.20	0.52	> 0.32	0.10	0.05	90.0	90.0	0.05	105.00	150.00	1.00 >	24.00	2.20	0.22	100.00	14.00	3.6	138.00	320.00	32.00 >	0.86	175.00	320.00	167.00	
	52 31	4.16.	0.32	0.32	0.00		15.90	0.23	1.07	0.05	9.0	6.73	0.10		0.01	0.0	0.05	0.0	3.05	100.00	0.53 >	1.03	0.61	0.87	. 32.00	2.00	3.	12.40	22.90	5.5	0.00	0.00	0.00	20.30	
	Diff.	0.739	- AH	¥	0.699	0.807 <	0.941	0.754	0.898	0.992	0.736	1.044	¥	· ≨	0.911	0.735	0.603	1.043	0.721	0.723	0.691	0.958	¥	\(\)	MA	0.731	0.363	0.881	0.866	0.488	0.00	0.586	0.758	0.589	
-	*	9 SO6	: 9	8	9 019	90 6		ALD 6	0 TE2			2XK	/	:	1000 6			SXZ O		9 SYF			: 9	:	/	804 6	200		SH6					N N N	
3	Date	04/19/89	10/03/86	06/01/88	05/10/89	01/24/89	01/18/90	05/10/89	01/18/90	02/07/89	08/23/89	11/01/90	12/21/87	03/01/88	02/01/89	08/23/89	02/13/90	11/01/90	03/15/89	12/14/89	08/23/89	03/06/90	11/14/86	05/18/88	08/28/87	03/28/89 POB	No/62/00	03/15/89	12/12/89	08/29/89	03/13/90	04/12/90	05/03/90	05/30/90	
1	sulp.	33	_	_	1,	94	8	5	3	9	7	~	2	7	9	7	3	7	25	25	28	82	5	\$	27	23	17	25	25	E C	\$ 8	65	\$!	19	
		0217 0217	2720	2220	2220	0303	0303	0347	0347	0360	0360	0360	0361	0361	0361	0361	0361	0361	9160	9160	1019	1019	1089	1089	1215	1215	CLZI	1217	1217	1337	1337	1337	1337	1557	

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Assay Type	H			ĒĒ	ĒĒ	FF		CPE	CPE MTT	CPE CPE	CPE CPE	CPE	CPE CPE
ŢĀ	10.11	32.44 50.83 9.90	19.28 7.15 2.47	12.01	13.43	40.04	32.53 4.32 7.03	1.60	0.00	2.70 0.00 5.70	1.70 0.00 14.44	1.50° 25.78 11.19	1.80
3	1.56 > 2.85 >	7.15 × 26.49 2.25	1.06 \$	3.63 v	1.07 >	30.00	7.50 \$ 0.00 \$ 0.40 \$	5.20 0.00 0.00	0.00	30.50 0.00 0.00 v	1.40 0.00 28.30	1.50 4.17 > 3.54	3.60
VI 98	0.00	12.56	0000	0.00	0.00	0.00	0.00	3.20	3.10 0.00 0.00	3.20	0.00	3.20 0.00 0.00	0.00
25 25	320.00	320.00 × 1000.00 × 811.00	320.00 320.00 320.00	320.00	320.00 320.00	320.00	100.00 100.00 1000.00	320.00 100.00 95.70	100.00 94.30 301.00	320.00 32.00 320.00	320.00 320.00 100.00	320.00 320.00 943.00	100.00 320.00
26 21	0.00	211.00 v 79.60 v 0.00	0.00	0.00	0.00	0.00	988	100.00 0.00 0.00	32.00 0.00 0.00	100.00 0.00 v	0.00	0.00 0.00 0.00	0.00
AI 50	1.56	7.15 37.82 3.52	51.30 0.00 0.00	9.00	1.07	36.00	10.00 0.00 2.63	5.20 - 0.00 0.00	7.00 - 0.00 0.00	30.50 ~ 0.00 0.00	1.41 0.00 38.50	1.50 ~ 5.54 6.45	3.60
TC 50	320.00 >	320.00 × 799.00 87.40	320.00 × 320.00 95.80	320.00 > 616.00	320.00 >	320.00 > 320.00 >	100.00 > 100.00 497.00	100.00 55.00 47.10	100.00 43.30 127.00	320.00 32.00 210.00	320.00 320.00 100.00	100.00 320.00 × 433.00	100.00 32.00
10 50	205.00 >	44.70 > 21.10 24.80	6.23 v 0.00 v	82.10 > 0.00	300.00 >	8.80 0.00	9.70 × 0.00 × 189.00	19.20 0.00 0.00	14.20 0.00 0.00	10.50 0.00 0.00	227.00 > 0.00 > 2.59 >	66.00 57.70 > 67.20	17.30 8.96
AI 25	3.40	15.38 40.13 3.55	2.24 0.08 3.80	1.39	1.85	86.00 0.00	88.00 0.00 1.77	0.00	27.80 0.00 0.00	0.00	58.20 0.00 45.60	2.10 6.94 5.56	31.30
70 25	320.00 >	320.00 > 559.00 55.90	6.60 3.20 21.50	298.00 146.00	320.00 >	260.00 61.00	73.00 74.50 74.50	66.00 27.30 25.10	100.00 11.40 49.00	320.00 16.00 86.40	320.00 320.00 73.50	66.00 241.00 238.00	16.00
S 21	94.20 >	20.80 v 13.90 15.80	2.94 42.50 5.66	47.50	173.00 >	3.00	0.80	6.20 0.00 100.00	3.60	2.80 0.00 0.00	5.50 × 0.00 × 1.61	32.00 34.70 42.70	3.20
Diff.	0.731	0.837 0.886 1.043	0.803 0.527 0.696	0.665	0.779	0.767 0.567	0.595	0.607 0.676	0.607 0.714	NA NA 0.870	NA NA 0.385	0.870 0.594	A A
=	38	SXS	M M M M M M M M M M M M M M M M M M M	8 00M	9 9C8	P0C 800	PF.3	S RF2	RF2	600	7	2 GC9	
Test	03/13/90 UJJ 05/03/90 WA	03/13/90 05/03/90 11/01/90	01/24/89 08/29/89 05/30/90	11/16/88	06/14/89 QCB 08/08/89 R2L	03/28/89 PDC 08/29/89 RDD	04/11/89 PFJ 08/29/89 RDE 05/31/90 WHF	10/09/86 08/30/89 RF2 05/31/90 WHG	10/09/86 08/30/89 RF2 05/31/90 WHH	10/09/86 08/21/87 06/14/89 qc9	10/17/86 10/29/87 05/16/89 q2T	10/21/86 06/14/89 QC9 08/08/89 R2M	11/07/86 06/01/88
Ship-	33	333	46 53 67	45	28	3 23	222	1 1 67	1 1 67	29	1 27	% % M	00
AVS S	1355	333	1654 1654 1654	55. 55.	1774	1841	1850 1850 1850	1973 1973 1973	1974 1974 1974	1975 1975 1975	1980 1980 1980	1992 1992 1992	2026

Table 18 (Cont'd)

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Assay	3 8	E	GPE	ËĒ	ě	i E		HTT	SPE CPE	9	CPE	HTT	MIT	H	CPE	CPE	H	ĔĔ	CPE	H	HTT	Ę,		CPE	H	HT	Ë				Ē		Ē	TH	
141	2.10	3.66	1.40	0.12	2 10	25. 25	7.0	5.57	8	0.30	0.60	0.00	8.00	12.52	3.30	2.90	65.81	40.27 50.58	4.40	2.76	4.86	13.73	<u>*</u>	1.80	52.20	1.7	86	40.18		6.93	3.94	77 21	6.25	0.00	
IS	1.50	0.00	1.40	^ 0.0	7 80	20.4	60.4	1.57	24.20	0.70	1.20	0.00	0.00	0.65 >	12.00	36.00	53.10 >	12.27 > 18.27 >	4.40			3.19 >		1.80	36.50 >	0.0	0.00	2.7		0.00	1.01	27 6	1.36	0.00	
A 98	0.00	0.00	0.00	0.0	5	3 23	6	0.00	0.32		1.00	0.00	0.00	0.00	3.20	0.0	19.10	12.09	320.00	0.00	0.0	8.0	3.0	0.00	0.0	0.0	8.8	30		8.6	88	5	. 6	0.00	
70 %	21.00	8.87	320.00	320.00	200 00	320.00	320.00	320.00	32.00	320.00 >	320.00 >	100.00	320.00	320.00	320.00 >	320.00	320.00 >	1000.00				8.8		320.00	315.00	313.00	268.00	320.00		280.00	278.00	20 80	32.00	31.10	
10.85	0.00 v	0.00	0.00	0.00	9	8 8	0.00	0.00	100.00	320.00 >	320.00 >	0.00	0.00	0.00	100.00	0.00	16.80 >	82.70 >	1.00 >	0.00	0.00	0.0	8	0.00	0.00	0.0	8.6	800		0.00	8 8	9	0.0	0.00	
AI 50	1.50 ~	0.00	1.40	0.0	7 80	3.7	7	2.41	24.00 ~	0.73	1.30	0.0	0.00	0.65	39.00 ~			25.32	438.00 ~	3.65	0.00	9.8	14.36	1.80	67.30	0.0	8.6	20.2		8.6	2.39	77 2	2.20	0.00	
TC 50	3.15	2.47	320.00	320.00	200 00	200.00	215.00	226.00	100.00	100.00	100.00	100.00	236.00	320.00 >	320.00 >	320.00 >	320.00 >	417.00	320.00	3.65 >	8.00	9.9	3.0	32.00	172.00	166.00	61.30	234.00		65.80	60.30	20 30	19.00	17.10	
10 50	0.00	0.00	236.30 >	0.00	41 00 4	28 70	30.60	93.80	4.10	134.00	80.00	0.00	0.00	< 00.96	8.20 >	8.90 >	5.27 >	17.90 28.60	0.73 >	1.00	0.00	0.67	0.0	18.00	2.56	0.00	0.6	22.50		9.00	22.50	6	8.61	0.00	
AI 25	2.10	0.00	3.40	8.8	9	28.50	7.16	5.69	100.00	0.32	9.1	0.00	91.0	11.70	31.00	100.00	78.50	32.98 46.68	10.00	1.50 <	3.7	8.09	8	10.00	67.80	3.09	00.0	167.07		2.61	1.3	72 2	2.89	0.00	
70 25	6.60	1.73	320.00	433.00	200 00	155 00	125.00	147.00	100.00	32.00	99.99	100.00	3.20	320.00 >	100.00	320.00	280.00	220.00 522.00	3.20	1.50 >	2.18	2.13	۲.۶۵	32.00	93.50	57.50	27.23	167.00 \$		45.30	25.50	14. 40	11.70	9.46	
10 25	3.20	0.0	95.00	0.0	90	7	17.50	24.80	1.00	100.00	99.99	0.00	19.60	27.40	- 3.20	~ 3.20	3.57	6.66	~ 0.32		0.58	0.26	7**0	3.20	1.38	18.60	9.6	1.00		7.07	14.70	78 2	4.07	0.00	
Diff.	NA 0.931	0.594	\$	0.841	1	272 0	0.581	0.653	¥.	¥	4	0.582	0.596	0.555	¥		0.701	0.722	¥		909.0	0.653	0.0	4	0.689	0.608	0.610	0.570		0.592	0.833	597 0	0.913	0.795	
14	: 5	RZH C	:	SNE					:	2	:	1 V9R	O XVO			1		SSY SW7	:	PFL C		33						XIA		Y KILY		310	13	O WI	
Test	11/05/86	08/08/89	11/07/86	11/30/89 05/31/90	12/18/84	04/11/80	08/30/89	05/31/90	02/01/86	08/21/87	09/21/87	04/12/90	05/06/90	07/31/90	03/02/87	08/21/87	04/11/89	12/05/89 12/12/89	12/30/86	04/11/89	08/30/89	05/31/90	01/16/7	12/31/86	04/11/89	08/30/89	04/12/90	07/12/90		08/30/89	05/03/90	11/28/80	03/13/90	05/03/9	
Ship-	o %	25	10	ıv v	5		12	29	Ξ	=	=	9	65	7	=	=	53	22	13	53	13	29	ò	13	53	T	65	6	;	£ \$	88	×	\$	\$	
AVS :	2034	2034	2138	2138	X25	1 %	2275	2275	2290	2290	2290	2290	2290	2290	2309	5309	2309	2309	2318	2318	2318	2318	0.00	2320	2320	2320	2320	2320		2362	2363	2570	2453	2453	

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Assay Type				FF	FF		EE	FFF	CPE	ĒĒ	S S S	S I I I I	
IAI	3.91 7.06 15.35 15.41 15.06	27.16 29.71 8.75	28.54 0.12 9.32 0.31 8.65	10.54	0.00	6.35	23.47	16.02 2.49 0.26	2.00 1.00 2.16	28.46	07.0	19.34 13.37 0.00	
5	1.82 × 4.46 × 2.13 5.46 × 2.49 ×	5.30 × 6.68 0.00	4.51 v 0.00 1.76 0.00 v	1.12 >	0.00	1.23 \$ 0.00	0.00 \$	3.91 >	3.70	00.00	10.00	4.60 2.87 v 3.05 v	
A1 %	% 88888 *	33.94	3.5. 0.00 0.00 0.00	0.00	0.00	0.00	0.00	0.00	3.10	0.00	3.20	0.00 0.00 0.00	
70 % 21	320.00 10.00 10.00 32.00	10.00 320.00 >	3.20 × 32.00 10.00 3.20 10.00	320.00 1000.00	320.00 1000.00	10.00 94.40 95.10	320.00	100.00 100.00 100.00	100.00 32.00 93.00	100.00	320.00 >	0.032 0.01 > 0.10 0.32	
S 21	98888	9.43 \$	0.00 0.00 0.00 0.00 0.00	0.00	0.00	0.00	0.00	0.00	32.00	0.00	100.00 >	0.00.0	
N 50	5.40 5.41 13.35 5.09	5.30 22.18 0.00	8.14 0.00 3.26 0.00 2.03	1.12	0.00	1.23	3.62	3.91	37.00 4.40 0.00	100.00	10.00	4.60 2.87 4.58 0.00	
TC 50	4.40 × 10.00 × 10.00 × 3.07	10.00 × 86.40 27.60	3.20 × 7.45 8.19 3.20 1.56	320.00 >	320.00	10.00 > 44.30 51.40	320.00 821.00	100.00 100.00 100.00	100.00 32.00 > 30.00	100.00 >	320.00	0.032 0.01 > 0.03 0.32	
10 50		1.89 × 3.89 0.00	0.39 × 0.00 2.51 0.00 × 0.77	255.00 >	0.00 > 235.00	8.14 v 0.00 0.00	0.00 > 227.00	25.50 > 0.00 > 0.00 >	27.00 ~ 7.20 > 0.00	1.00	32.00 118.00	0.00 > 0.00 > 0.01	
AI 25	1.82 < 3.29 8.35 4.98	8.24 12.57 4.70	8.50 0.00 2.81 0.00 3.54	1.98	0.00	1.96	130.48	0.00	10.00 3.10 0.40	11200	5.00	10.00 5.31 4.83 0.00	
TC 25	1.82 × 1.62 4.09	10.00 > 26.00 8.74	5.33 5.33 3.20 0.73	320.00 >	273.00	10.00 × 22.40 27.40	283.00 374.00	100.00 × 100.00 51.00	10.00	67.70	100.00	0.03 0.01 > 0.02 0.32	
10 25	1.00 0.45 0.49 0.30	1.21 × 2.07 1.86	0.21 0.00 1.57 0.30 v	161.00 >	0.00	5.10 v 0.00 5.72	2.17	10.00 0.00 0.00	3.20	0.00	21.00	0.003	
Diff.	0.689 < 0.503 0.791 0.704 1.051	0.480	1.002 0.724 0.491 0.733 1.116	0.531	0.531	0.394 0.570 0.936	0.506	0.791	NA - 0.976	0.463	¥ ¥	NA - 0.837 0.814 0.912	
÷ *	39 PFH 39 SLF 30 XGZ 30 XGZ	39 SLG 20 UJK 20 VVT	39 OKN 39 952 39 SLH 39 SSH 30 ZXU	39 QK3 90 V9T	39 QK3 90 1PY	39 SLJ 30 V9U 30 1PY	39 SLL 39 SLL	38 088 39 SLM 90 190	39 OES	39 PO1 90 VE9	37	39 OKR	
Test Oate	04/11/89 11/28/89 05/31/90 07/12/90	11/28/89 03/13/90 05/03/90	02/07/89 05/23/89 11/28/89 12/05/89	06/27/89 QK3 04/12/90 V9T	06/27/89	11/28/89 04/12/90 12/05/90	11/28/89 SLL 04/12/90 V9V	12/21/88 11/28/89 12/05/90	05/27/89 05/25/88 01/24/89 0ES	04/21/89 PO1 04/17/90 VE9	04/28/87 08/21/87	03/01/88 02/07/89 12/06/89 12/05/90	
Ship- ment	53 67 67	222	48 15 21 48 15/21	57 65	57	65 67	91	555	223	25 65		32 61 61	
AVS.	2503 2503 2503 2503 2503	2506 2506 2506	2563 2563 2563 2563 2563	2573 2573	2575 2575	2580 2580 2580	2590	2604 2604 2604	2716 2716 2716	2720	2743	2812 2812 2812 2812 2812	

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Assay Type	H	ĒĒ	EEE	9 CP E		EE	ËË	ĦĦ	ĔĔ	ĒĒ	ĔĒ	###	ËË	H
TAI	21.14	24.44	3.15	2.80	23.21 11.88 24.37	1.93	7.45	15.85	10.82	41.87	2.66	0.00 37.49 14.64	0.62	3.04
18	2.51	6.50 >	0.00	10.00	4.10 v 5.29	1.18	0.00	1.71	1.69 \$	12.20	1.65	0.00	0.00	0.00
VI 95	3.39	0.00	0.00	3.20		0.00	0.00	3.22	0.00	9.85	0.00	0.00	0.00	0.00
70 95	320.00	320.00	27.40 92.60 100.00	32.00	3.20	100.00	94.70	95.80	320.00	96.40 100.00	31.20	3.20 320.00 >	100.00	100.00
10 %	94.50 × 0.00 ×	0.00	0.00 9.28 0.00	3.20	0.00	0.00	0.00	29.70	0.00	0.00	0.00	29.00 >	0.00	0.00
AI 50	4.61	10.38	0.00 0.00	-	3.43	1.18	1.54	3.18	1.69	16.90 0.00	2.49	29.80	3.0	0.00
TC 50	261.00	210.00	3.82 9.62 13.50	32.00	3.20 4	100.00 > 657.00	34.60	48.70	320.00	64.20	19.40	3.20 229.00 210.00	32.00 63.50	30.80
10 50	56.60	20.20	0.00 0.00	0.30	0.78 > 0.59	84.40 >	39.50	15.30	190.00 >	3.73	7.79	0.00 × 7.68 0.00	0.00	0.00
AI 25	3.33	0.00	1.11	10.00	3.23 12.22	2.17	1.52	2.47	3.62	23.20	2.58	0.00 0.46 8.22	0.00	0.00
70 25	142.00	131.00	1.75 6.41 6.39	3.20 >	3.20 >	100.00 > 485.00	25.30	26.20	320.00 > 62.20	46.30	12.90	3.20 2.10 32.00	10.80	21.00
10 25	42.50	13.40	1.57 3.25 8.72	0.32		46.00 \$	16.60	10.60	88.50 >	2.00	4.99	0.00 × 4.62 3.89	0.00	0.00
Diff.	0.593	0.546	0.729 0.515 0.572	¥ 4 4	586 532 832	0.993	0.515	0.716	0.732	0.744	0.704	0.555 0.711 0.783	0.614	0.614
Test Pit Date #	04/11/89 PFN 09/07/89 RIR	09/07/89 RIS 03/13/90 UJN	02/08/89 OLU 04/19/89 PHT 09/07/89 RIU	03/01/88		12/20/88 07N 04/17/90 VEF	04/19/89 PHT 04/21/89 PAA	04/19/89 PMX 04/21/89 PAE	04/17/90 VEJ 07/06/90 XAS	04/19/89 PN6 12/20/89 T0B	04/05/89 PC4 04/19/89 PHY	11/03/88 0CU 04/19/89 PN1 01/18/90 TF3	11/03/88 OCV 04/19/89 PH2	11/03/88 OCV 04/19/89 PN2
Ship- Te ment Da	53 04/ 26 09/	26 09/	48 02/ 27 04/ 25 09/	25 12/ 25 03/ 25 03/		31 12/	32 04/	32 04/	65 04/	33 04/ 33 12/	32 04/	32 11/ 32 04/ 32 01/	32 11/	32 11/
AVS Sh No. me	2906 5	2907 2	2979 2	2980 2 2980 2 2980 2		3374 3	3425 3	3450 3	3491 6	3494 3	3565 3	3586 3586 3586 3	3588 3	3589 3

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Assay Type					A T T T		FEF		CPE	OPE HTT
IA	19.63 35.74 4.38 19.84	1.35	21.94 30.31 4.07 59.40	4.61 22.75 14.28 22.98	1.85° 14.01 13.05	22.39 9.98 8.61	6.49 3.55 1.98	9.82 0.00 3.36 6.39	1.40	1.50
S	1.24 × 0.00 × 3.53 × 3.39 ×	0.00 0	4.66 9.08 v 42.71 v	0.00 4.06 > 3.13 > 4.58 >	2.10 2.11 > 2.61	4.35 > 1.95 2.09	0.00 > 0.00	2.56 v 0.00 v 0.00 v 1.08 v	3.70	2.00
AI 95	0.00 0.00 0.00 0.00 0.00	0.00	0.00	0.00	3.20 1.08 > 0.00	3.13	0.00	0.00	3.10	3.10
70 95	328.00 320.00 × 320.00 320.00	320.00 × 1000.00 1000.00	953.00 10.00 10.00 10.00	320.00 320.00 148.00 32.00	32.00 3.20 > 80.60	1000.00 1000.00	0.03 26.50 9.66	320.00 1000.00 2.85 1.00 1.00	31.00	100.00 87.00
10 95	0.00 0.00 0.00 0.00 0.00 0.00	^ ^ ^	00.00	00.00	10.00 2.97 > 0.00	320.00 > 0.00 > 0.00 >	0.00	000 000 000 000 000 000 000 000	32.00	32.00
A1 50	2.19 11.60 0.00 3.53 3.39	0.00	8.33 53.83 33.22 42.71	0.00 6.97 6.39 9.16	21.00 2.11 5.06	6.34 2.87 3.00	0.00	3.27 0.00 0.00 0.00 2.37	12.00 3.00	5.80
10 50	184.00 320.00 > 320.00 > 320.00 >	320.00 × 1000.00 1000.00	528.00 10.00 v 10.00 v	153.00 188.00 77.30 24.50	32.00 ~ 3.20 > 12.40	860.00 715.00 656.00	0.03 2.71 > 1.16	320.00 × 344.00 1.00 1.00 0.61	100.00 ~	100.00 ~
10 50	83.90 27.70 > 0.00 > 283.00 >		63.40 0.19 > 0.30 > 0.23 >	0.00 27.00 12.10 2.68	1.50 ~ 1.52 > 2.46	136.00 249.00 218.00	0.00	97.80 × 0.00 × 0.00 × 0.00 × 0.26	8.50 ~	4.90 ~ 4.00
AI 25	4.80 22.10 0.00 21.43 7.04	0.00	8.78 13.25 1.89 65.93	1.50 9.54 5.36 22.60	10.00 3.06 6.71	8.40 3.45 3.29	4.37 1.69 < 0.00	4.47 0.00 1.00 2.16	10.00	3.10
10 25	104.00 189.00 320.00 1000.0 >	320.00 × 1000.0 1000.0	296.00 1.69 0.27 10.00 >	77.90 109.00 37.80 12.30	3.20 > 6.42	590.00 485.00 456.00	0.03 > 1.69 > 0.67	250.00 91.50 1.00 ~	32.00 ~	10.00 -
52 21	21.70 8.55 0.00 \$ 46.70 \$	67.50 \$ 0.00 \$ 0.00 \$	33.70 0.13 0.14 0.15 v	51.90 11.50 7.06 0.54	0.32 1.05 × 0.96	70.20 141.00 139.00	0.00	55.90 0.00 1.00 × 0.13	3.20	3.20
Diff.	1.001 0.859 0.677 0.814 0.751	0.634 0.802 0.944	0.641 0.846 0.704	0.754 0.594 0.887 0.706	NA 0.678 0.723	0.669 0.817 0.859	0.718 0.894 < 0.699	0.721 0.565 0.783 < 0.628 0.598	NA ~ 0.615	NA ~ 0.976
P1.	789 OA6 789 DUV 789 SLR 789 SQB 790 WS9	789 SSX 790 ZVV	89 51.8 790 WHM	05/15/90 W5A 07/17/90 XL9 03/01/89 OUR 03/22/89 PBD	88 89 OLV	789 SSR 790 TF2 790 TVD	789 OLY 789 SUE 790 UOD	90 WSE 90 XCU 90 TF3 90 U07	05/11/88 04/11/89 PF0	05/11/88 01/24/89 0ES
Test	E 01/12/89 03/07/89 11/28/89 12/06/89 05/15/90	11/03/88 12/05/89 10/30/90	12/20/89 11/28/89 05/31/90 07/12/90	05/15/90 07/17/90 03/01/89 03/22/89	04/12/88 02/08/89 12/07/89	12/05/89 01/18/90 02/08/90	02/08/89 12/07/89 02/13/90	05/15/90 07/06/90 01/18/90 02/06/90 02/13/90	05/11/88	05/11/88
Ship- ment	ABEBE 51 51 65		12 x22	88 BB	28 89 48 89	888	8 8 8	88 888	23 39	6 9 6 9
AVS No.	3592 3592 3592 3592 3592	3612 3612 3612 3612	3621 3802 3802 3802	3819 3864 3964	0207 0207 0207	4071 4071 4071	7207 7207 7207	4075 4075 4223 4223 4223	4240	4241

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Assay Type	3 8			FFF	ĒĒ		FFFF	FFFF		ĒĒ	
Ĭ	1.30	7.0¢ 5.41 0.00	15.70 12.66 7.32	5.31 12.39 2.65	17.09	15.02 1.03 0.00 8.93	8.63 2.20 14.26 12.80 8.44	24.18 0.89 19.51 9.83 4.95	0.45 9.97 28.27 12.35	5.88	0.00 0.00 21.56 6.89
15	0.00	1.27 > 0.00	0.00 × 2.51 0.00	2.93	3.58	2.31 \$ 0.00 \$ 2.03	1.07 v 0.00 v 2.46 2.23 v 2.07	3.17 0.00 × 4.56 1.81 0.00 ×	1.00 × 0.00 × 5.22 × 2.21	0.12 >	0.00
A1 95	0.00	988	0.00	0000	3.20	9888	00000	3.37	0.00 0.00 10.55 0.00	0.00	0.000
70 95	320.00 >	10.00 87.90 85.20	32.00 100.00 94.60	10.00 32.00 32.00	320.00 >	90.60 100.00 100.00 94.30	26.80 10.00 9.13 10.00	100.00 100.00 100.00 100.00 57.50	3.01 0.10 1.00 × 0.89	320.00 953.00	94.60 95.90 144.00 95.50
10 95	>320.00 >	0.00 0.00	0.00	0.00	100.00 >	0.000	000000	29.70 0.00 0.00 0.00 0.00	0.0000	0.00	0.000
A1 50	0.00	1.27	0.00 5.30 0.00	0.00 5.11 0.00	5.40	4.92 0.00 0.00 2.98	2.34 0.00 4.91 3.55 3.00	5.82 0.00 3.83 0.00	1.88 0.00 10.02 3.54	2.41	0.00 0.00 7.73 2.19
TC 50	0.32	10.00 × 25.30 × 22.40	28.70 51.60 25.30	9.73 12.00 8.27	308.00	24.70 100.00 100.00 27.70	5.98 5.82 2.62 8.18 8.43	88.40 59.60 83.80 77.80	0.60 v 0.10 0.43 0.25	320.00 > 278.00	46.00 50.30 61.10 54.70
10 50	320.00 <	7.90 v 0.00 0.00	0.00 0.00	0.00 2.35 0.00	56.90	5.01 0.00 9.30	2.55 0.00 0.53 2.30 2.81	15.20 0.00 9.50 24.90 0.00	0.32	133.00 >	0.00 0.00 7.89 25.00
AI 25	0.10	1.99	4.38	2.72 4.62 1.74	3.85	6.47 0.00 3.72	2.39 0.00 5.30 4.20	4.60 0.00 3.98 0.00	1.00 < 2.27 11.43 3.90	1.13	39.06
TC 25	0.32	10.00 v 17.60 13.50	18.30 24.50 17.60	6.40	204.00	11.60 100.00 91.50 18.90	2.72 2.57 1.31 5.14 5.82	48.10 22.60 43.30 45.10 7.06	0.32 - 0.10 > 0.22 0.16	16.50	17.30 21.70 39.10 >
10 25	3.20	5.03 × 6.52 0.00	4.17 5.18 5.73	2.35 1.49 3.20	41.60	0.00	1.14 0.00 0.25 1.22 1.64	10.50 0.00 4.97 11.30 0.00	0.32	14.60	0.00
oiff.	\$ \$	0.340	0.505 0.787 0.885	0.385 0.787 1.037	0.691	1.121 0.635 0.648 0.588	0.998 0.353 0.855 0.635 0.588	0.968 0.353 0.855 0.701 0.615	0.979 < 0.979 0.722 0.603	0.725	0.713 0.768 0.676 < 0.779
= =	88/ 88/	/89 SH3 /90 W5J	/89 SH4 /89 SG9 /90 ZXU	/89 SMS /89 SQ9 /90 2XX	/89 QP1 /89 R81	/89 0JN /90 TRN /90 UOE /90 UOS	/89 OUO /89 SA7 /90 TRN /90 UOB	/89 SUA /89 SUA /90 TRO /90 U09	/89 OUG /89 SSY /90 UOA	/90 WSL	/89 GP2 /89 R81 /90 UEU /90 VAU
Test Oate	04/13/88	11/28/89 05/15/90 07/06/90	11/28/89 12/06/89 11/01/90	11/28/89 12/06/89 11/01/90	07/07/89 08/16/89	02/01/89 02/06/90 02/13/90 02/13/90	03/01/89 11/28/89 12/06/89 02/06/90 02/13/90	03/01/89 11/28/89 12/06/89 02/06/90 02/13/90	03/01/89 03/01/89 12/05/89 02/13/90	05/15/90 07/06/90	07/07/89 08/16/89 03/08/90 05/03/90
Ship-		2 5 5 5	255	255	33	77.75	63 64 68 63 63 63 63 63 63 63 63 63 63 63 63 63	48 63 63	84 4 8 8 8 8 8 8 8 9 8 9 8 9 9 9 9 9 9 9	65	3333
AVS	4275	4277 4277 4277	4278 4278 4278	4281 4281 4281	4452	4527 4527 4527 4527	4590 4590 4590 4590 4590	4592 4592 4592 4592 4592	6097	4611	433 433 433 433

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Assay Type	T I	EE	EE	ĔĒ	FFF	ĒĒ	FFF	ĒĒ	FFF	FFF	EEE	ĒĒ	ĒĒ	EE	
¥	10.13	10.27	12.63	12.94	23.97 13.78 14.58	13.19	30.97 22.50 32.83	20.27	34.34 18.49	12.8 0.00 2.8	43.84 2.21 13.52	3.72	9.93	14.30	
13	0.18	0.00	2.20	2.37	7.19 > 0.00 > 2.68	2.16	13.80 > 5.26 > 10.03	3.53 >	8.12 3.49 0.00	1.57	25.10 > 0.00 > 2.89 >	32.00 > 0.00 >	2.00 >	1.98 •	
AI 95	3.10	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00 10.28 0.00	0.00	0.00	0.00	1.07 > 0.00	1.07 >	
70 95	310.00	309.00	307.00	308.00	320.00 32.00 32.00	320.00	320.00 100.00 320.00 >	320.00 320.00	96.70 320.00 > 96.60	320.00 320.00 1000.00	320.00 320.00 1000.00	320.00 320.00	320.00 >	320.00 >	
S6 21	0.00	0.00	0.0	0.00	0.00	0.00	0.00 0.	0.00	31.10 >	0.00	0.00	0.00	299.00 > 0.00 >	299.00 \$	
AI 50	3.56	2.80	3.30	3.35	29.10 0.00 8.08	3.99	46.60 5.26 19.28	7.07	12.50 5.04 0.00	3.51 0.00 0.00	25.10 0.00 3.47	32.00	2.00	1.98	
1C 50	202.00	187.00	192.00	202.00	29.10 × 11.80 12.10	275.00	73.60 100.00 v 273.00	320.00	59.40 83.50 66.00	240.00 227.00 188.00	320.00 > 320.00 1000.00 >	320.00 >	320.00 >	320.00 >	
10 50	0.00	0.00	58.20	0.00	1.00	69.00	1.58 19.00 > 14.20	42.50	4.76 16.60 0.00	68.40 0.00 0.00	12.70 > 0.00 > 289.00 >	10.00 \$	160.00 > 0.00	161.00 > 0.00	
AI 25	0.24	3.23	3.57	3.67	7.19 < 9.10 5.73	4.02	21.70 7.40 17.99	10.40	13.20 4.95 2.52	2.52 0.00 3.13	72.20 0.00 5.52	140.00	2.83	0.00	
TC 25	10.00	119.00	128.00	143.00	7.19 × 4.37	149.00	21.70 > 100.00 > 142.00	150.00 181.00	38.60 57.80 49.00	107.00 134.00 71.80	320.00 > 320.00 835.00	320.00 >	320.00 > 88.70	320.00 > 524.00	
10 25	41.20	36.90	35.90	38.80	1.00	37.10 55.30	13.50 > 7.89	14.40	2.94 11.70 19.40	42.50 0.00 23.00	4.43 > 0.00 > 151.00	2.28 > 0.00 >	113.00 >	115.00 > 0.00	
Diff.	0.511	0.505	0.541	0.541	0.929 < 0.456 1.037	0.885	0.885 < 0.481 0.922	1.026	1.240 0.403 0.791	0.475	0.337 0.340 0.791	0.409	0.377	0.531	
¥ *	11/08/88 031 02/13/90 U0F	/88 032 /90 U0G	788 033 790 UOG	788 033 790 UCB	89 09N 89 SN9 90 ZXX	89 095 89 SMA	89 098 89 SMA 89 SQB	89 OCO	89 OJS 89 SMB 89 SYE	89 OHR 89 SNC 89 SQB	89 OHS 89 SHC 89 SQC	89 OHT	789 OPG	02/21/89 OPH 02/13/90 UOH	
Test Date	11/08/88 02/13/90	11/08/88 02/13/90	11/08/88 02/13/90	11/08/88 03/06/90	01/12/89 11/28/89 11/01/90	01/12/89	01/12/89 11/28/89 12/06/89	01/17/89	02/01/89 11/28/89 12/14/89	02/14/89 11/28/89 12/06/89	02/14/89 11/28/89 12/06/89	02/14/89	02/21/89	02/21/	
Ship-	33	22	22	23	333	99	332	99	8 8 8 8 7	84 87 97	87 87 91	84 84	84	87	
AVS Ho.	4765	4768	6927	7120	4785 4785 4785	4795	9627 9627 9627	4809	4822 4822 4822	4825 4825 4825	4827 4827 4827	4829	4843	7787	

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Assay	FFF	FFF	FF		FF	FFFF	FFF	EE	FFF	THE LE	ĔĒ	ĔĒ	EEE	T I
ĭ	64.54 1.24 0.73	3.17	5.31	53.33 17.29 0.00	13.74 2.36	60.86 41.95 33.73	50.93 6.17 6.60	50.97	2.69	21.31	13.61 20.53	0.87	27.87 5.06 15.05	2.46
5	104.00 v	239.00 > 0.00 > 0.00	58.30 >	217.00 × 0.00 × 0.00	2.52 > 1.04 >	20.50 > 11.28 > 20.93 > 8.41 >	47.10 > 0.00 > 0.02 >	129.00 >	0.00 0.34 0.01	4.23	0.00 >	0.00 >	5.27	0.00
N 95	4.47 × 0.00 0.00	988	0.00	888	0.00	19.50 10.85 5.66 >	989	11.50	0.00	0.00	0.00	0.0	0000	0.00
70 %	320.00 × 320.00 1000.00	320.00 320.00 1000.00	320.00	320.00 320.00 1000.00	29.30	328.00 320.00 v 320.00 v	320.00 320.00 1000.00	320.00 >	320.00 320.00 1000.00	320.00	320.00 1000.00	9.39	312.00 307.00 304.00	335.00
10 %	71.50 > 0.00 > 0.00 >	0.00	0.00	0.00	0.00	16.80 29.50 × 56.60 × 142.00 ×	0.00	27.80 > 0.00 >	0.00	0.00 > 239.00 >	0.00	0.0	0.00	0.00
A1 50	0.00	0.00	106.00	. 217.00 0.00 0.00	4.46	36.20 15.47 20.93 8.41	47.10 0.00 2.08	235.00	0.00	7.17	0.00	0.00	9.81	3.56
TC 50	320.00 × 320.00 1000.00	320.00 3 320.00 1000.00	243.00	320.00 × 320.00 1000.00	10.00	184.00 218.00 320.00 >	320.00 × 320.00 1000.00 ×	320.00	320.00 320.00 > 259.00	320.00 >	320.00	3.93	170.00 194.00 161.00	185.00 189.00
10 50	3.8 0.0 0.8 0.0	1.3¢ 0.00 0.00 0.00	2.30	1.47 \ 0.00 \ 0.00 \	1.73	5.08 14.10 15.30 v	6.80 v 0.00 v 481.00 v	1.09	0.00 × 267.00 × 978.00	44.60 >	0.00 \$	0.00	17.30 97.30 21.70	0.00 53.10
AI 25	250.00	. 320.00 0.12 0.00	134.00	320.00 5.66 0.00	4.36	104.00 18.62 35.67	320.00 2.68 0.06	7.65	0.93 0.65 0.55	8.91	5.66	1.76	8.77 2.64 1.28	0.00
10.25	320.00 > 320.00 > 3.20	9.24 9.24 1000.0	134.00 >	320.00 > 100.00	4.36 >	104.00 v 159.00 v 320.00 v	320.00 > 320.00 > 8.64	140.00 >	274.00 90.90 13.70	189.00	320.00 > 738.00	1.76 > 0.50	91.40	102.00
10 25	1.28 > 0.00 > 283.00	< 1.00 × 74.30 × 0.00 ×	0.00	, 1.00 , 56.60 , 0.00 , 0.00	< 1.00 5.53 >	4 1.00 8.55 3.00 v	< 1.00 > 120.00 > 143.00	32.00	294.00 140.00 24.90	21.20	56.60 × 35.20	1.000.12	10.40	30.50
01ff.	0.429	0.435	0.435	0.591	0.454	0.470	0.401 0.858 0.853	0.405	0.930	0.904	1.354 0.634	1.267	0.820	1.003
st Pit	02/22/89 OPT 11/30/89 SWN 12/06/89 SQC	02/22/89 OPU 11/30/89 SWN 12/06/89 SQD	02/22/89 OPU 11/30/89 SNO	02/22/89 OPV 11/30/89 SNO 12/06/89 SQD	02/22/89 OPU 11/30/89 SNP	02/22/89 OPX 11/28/89 SMD 12/06/89 SQE 12/05/90 102	02/22/89 OP2 11/30/89 SNP 12/06/89 SQE	02/22/89 000 11/28/89 SME	01/17/89 OC2 05/15/90 W5M 07/06/90 XCV	01/17/89 OC3 12/06/89 SQF	01/19/89 006 12/20/89 10E	01/19/89 00E 12/20/89 10E	01/31/89 OHT 11/30/89 SNQ 12/06/89 SQF	02/01/89 0JH 12/06/89 SQG
Ship- Test ment Date	48 02/2 48 11/2 61 12/1	48 02/2 48 11/2 61 12/0	48 02/ 48 11/	48 02/2 48 11/2 61 12/4	48 02/ 48 11/	48 02/2 48 11/2 61 12/4	48 02/4 48 11/2 61 12/4	48 02/ 48 11/	46 01/ 65 05/ 65 07/	46 01/ 61 12/	46 01/ 46 12/	46 01/ 46 12/	46 01/2 46 11/7 61 12/9	46 02/1 61 12/1
AVS Sh No. me	9 8787	9 6787 9 6787 9 6787	7 0587	4851 4 4851 4 4851 6	4853 4	4855 4855 4855 4855 4855 4855	4861 4 4861 4 4861 6	7 5987	1984 1984 1984	4871 4	4875 4	4891 4	7 2687 7 2687 9 2687	4919 4

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FFF	EEE	FE	EE	E E	EE	ĔĔ	FE	EE	EEEE	EE	EEE		FF
22.59 1.92 0.00	24.44 1.42 0.00	23.05	3.27	24.04	20.36	4.08	15.77	14.70	19.13 5.37 39.83 4.08	27.34 6.52	39.13 9.89 4.93	42.55 19.28 15.38 27.23 10.11	22.52 9.08
4.00.0 0.00 0.00	2.49 0.00 0.00	1.93 >	0.00	3.65 >	20.60	0.00	1.12	1.35	5.20 × 0.00 14.55 0.00 ×	36.80	9.88 0.00 0.00 0.00	12.00 > 2.41 > 2.21 > 7.48 > 2.00 >	1.22 > 0.00
0.00	9.25	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00 11.95 0.00	0.00	28.50 0.00 0.00	0.00 0.00 0.00 0.00 0.00 0.00	0.00
83.00 32.00 32.00	84.50 100.00 30.80	320.00 289.00	32.00	320.00 1000.00	303.00	320.00 964.00	265.00	90.30	320.00 1000.00 1000.00 > 3200.00	320.00	285.00 100.00 296.00	320.00 × 320.00 × 1000.00 1000.00	320.00 1000.00
888	9.14 0.00 0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	83.70 0.00 0.00 0.00	0.00	0.00	320.00 × 293.00 × 0.00 × 0.00 × 0.00 ×	0.00
0.00	5.05 0.00 0.00	3.68	12.40	0.00	48.70	3.19	3.09	3.23	\$.20 0.00 31.62 0.00	36.80	0.00	12.00 2.41 2.21 7.48 2.00	1.22
20.50 20.00 19.90	20.40 100.00 17.00	252.00 85.10	22.20	320.00 864.00	149.00	320.00 638.00	18.60	21.50 28.60	320.00 × 1000.00 660.00	320.00 > 320.00	75.20 100.00 93.10	320.00 × 320.00 × 1000.00 × 1000.00 × 1000.00 ×	320.00 × 1000.00
0.00	4.05 0.00 v	00.0	6.00	0.00 >	3.05	200.00	6.03	%.0 %.0	61.60 v 0.00 v 20.90 v 0.00 v	8.69 >	4.31 0.00 >	26.60 × 133.00 × 453.00 × 134.00 × 501.00 ×	262.00 > 0.00 >
9.63 0.00 0.00	6.58 0.00 0.00	13.20	0.00	0.00	37.80	3.42	3.81	0.00	8.81 3.17 26.34 0.00	3.44	3.44 0.00	55.10 4.41 5.59 21.37 3.76	9.50
10.90 13.10 11.90	10.10 100.00 9.15	132.00 58.60	11.60 >	320.00	62.90 155.00	181.00	6.72	9.00	320.00 > 650.00 > 304.00	320.00 > 267.00	42.60 100.00 > 58.40	320.00 × 1000.0 × 1000.0 × 1000.0 ×	320.00 > 715.00
0.00	0.00	0.00	0.00	30.40	35.60	320.00 134.00	1.76	2.03	36.30 × 205.00 11.50 0.00 ×	5.27 >	29.00 >	5.80 × 72.60 × 179.00 × 46.80 × 266.00 ×	6.84 v
0.828 0.577 0.844	0.912	0.594	0.594	0.660	0.680	0.746	0.838	0.702	1.066 0.675 0.627 0.786	0.524	0.452 0.592 0.831	0.548 0.592 0.663 0.627 0.826	0.559
89 SWH 89 SWH 89 SY7	89 OUZ 89 SMI 89 SY7	89 OX1 89 SMI	89 OX1 89 SHJ	89 P1X 89 SSZ	89 P20 89 SML	89 P24 89 SQG	89 P25 89 SMN	89 OT1 89 SMO	89 0VA 89 SSS 89 TOF 90 TVE	89 014 89 SMO	89 OT6 89 SMP 90 TF4	89 SHP 89 SNP 89 SST 89 TOF 90 TVF	89 018 89 SU8
03/07/ 11/28/ 12/14/	03/07/ 11/28/ 12/14/	03/07/ 11/28/	03/07/	03/14/	03/14/	03/14/	03/14/	02/28/		02/28/	02/28/ 11/28/ 01/18/	02/28/ 11/28/ 12/05/ 12/20/ 02/08/	02/28/89 12/07/89
222	222	22	22	51 27	5 5	22	51	84	45 62 45 GABSN	8 8	87 87	88686	8 29
767	6£67 6£67	4942	4943	4978	5867	7665	5667	5035 5035	5040 5040 5040 5040	5048 5048	5053 5053 5053	5058 5058 5058 5058	\$060 \$060
	51 03/07/69 CMM 0.828 1.13 10.90 9.63 2.43 20.50 8.43 0.00 83.00 0.00 4.48 > 22.59 51 11/28/89 SMH 0.577 0.00 13.10 0.00 0.00 20.00 0.00 0.00 32.00 0.00 0	\$1 03/07/69 OMX 0.828 1.13 10.90 9.63 2.43 20.50 8.43 0.00 83.00 0.00 4.48 > 22.59 51 11/28/69 SMH 0.577 0.00 13.10 0.00 0.00 20.00 0.00 0.00 30.00 0.00 0	51 03/07/89 OMR 0.826 1.13 10.90 9.63 2.43 20.50 8.43 0.00 83.00 0.00 4.48 > 22.59 51 11/28/89 SMH 0.577 0.00 13.10 0.00 0.00 20.00 0.00 0.00 32.00 0.00 1.92 51 12/14/89 SMT 0.844 0.00 11.90 0.00 19.90 0.00 1.42 24.44 1.42 1.42 1.42 1.42 1.42 1.42 1.42 1.42 1.42 1.42 1.42 1.44	\$1 03/07/89 QMX 0.828 1.13 10.90 9.63 2.43 20.50 8.43 0.00 83.00 0.00 4.48 > 22.59 \$1 11/28/89 SMH 0.577 0.00 13.10 0.00 0.00 20.00 0.00 32.00 0.00 1.92 \$1 12/14/89 SY7 0.844 0.00 11.90 0.00 19.90 0.00	\$1 11/28/89 SWH 0.577 0.00 11.90 9.63 2.43 20.50 8.43 0.00 83.00 0.00 4.48 > 22.59 51 11/28/89 SWH 0.577 0.00 11.90 0.00 0.00 0.00 0.00 0.00 0.0	51 03/07/89 OMZ 0.828 1.13 10.90 9.63 2.43 20.50 8.43 0.00 83.00 0.00 4.48 > 22.59 51 11/28/89 SMH 0.577 0.00 13.10 0.00 0.00 20.00 <	\$1 17.28/89 \$WH 0.577 0.00 113.10 9.643 2.43 20.50 8.43 0.00 83.00 0.00 4.449 22.59 51 17.28/89 \$WH 0.577 0.00 113.10 0.00 0.000 20.00 0.00 0.00 0.00 32.00 0.00 0	51 11/28/89 SMH 0.577 11.13 10.90 9.63 2.43 20.50 6.00 83.00 0.00 4.48 22.59 51 11/28/89 SMH 0.577 0.00 11.90 0.00 20.00 0.00 52.00 0.00 32.00 0.00 1.92 51 11/28/89 SMH 0.577 0.00 11.90 0.00 20.00 0.00 52.00 0.00 0.00 1.92 0.00	11/28/99 SMH 0.577 0.00 15.10 0.00 0.00 20.00 0.00 23.00 0.00 0.00 1.92 11/28/99 SMH 0.577 0.00 15.10 0.00 0.00 0.00 19.90 0.00 0.00 23.00 0.00 0.00 1.92 11/28/99 SMH 0.577 0.00 11.90 0.00 0.00 0.00 19.90 0.00 0.00 23.00 0.00 0.00 1.92 11/28/99 SMH 0.577 0.00 11.90 0.00 0.00 0.00 19.90 0.00 0.00 20.00 0.00 0.00 0.00 11/28/99 SMH 0.577 0.00 12.00 0.00 0.00 0.00 0.00 0.00 0.00 0.00 0.00 11/28/99 SMH 0.502 0.00 9.15 0.00 0.00 0.00 0.00 0.00 0.00 0.00 0.00 11/28/99 SMH 0.554 0.00 12.00 0.00 0.00 0.00 0.00 0.00 0.00 0.00 0.00 0.00 11/28/99 SMH 0.554 0.00 12.00 0.00 0.00 0.00 0.00 0.00 0.00 0.00 0.00 0.00 0.00 11/28/99 SMH 0.554 0.00 12.00 0.00	51 11/28/99 SMR 0.527 1.13 10.50 9.63 2.43 20.50 6.00 52.00 0.00 52.00 0.00 52.00 0.00 52.00 0.00 52.00 0.00 52.00 0.00 1.22 9.14 9.00 1.00 1.00 1.00 1.00 1.00 1.00 1.00 1.00 0.00 1.00 0.00 1.00 0.00 1.00 0.00 0.00 1.00 0.00 </th <th>111/126/99 SMI 0.557 0.544 0.00 11.00 0.00 0.00 19.90 0.00 0.00 0.</th> <th>11 11 11 11 10 9.43 2.43 20.59 6.43 0.00 32.00 0.00 32.00 0.00 32.00 0.00 32.00 0.00 1.99 0.00 0.00 22.59 0.00 0.0</th> <th> 1,124/99 344 3.77 3.04 3.14 3.04 3.24 3.05 3.45 3.05 3.05 3.00</th>	111/126/99 SMI 0.557 0.544 0.00 11.00 0.00 0.00 19.90 0.00 0.00 0.	11 11 11 11 10 9.43 2.43 20.59 6.43 0.00 32.00 0.00 32.00 0.00 32.00 0.00 32.00 0.00 1.99 0.00 0.00 22.59 0.00 0.0	1,124/99 344 3.77 3.04 3.14 3.04 3.24 3.05 3.45 3.05 3.05 3.00

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FFF	E E	ĦĦ	###		FFF	ĒĒ	FF	ĒĒ	EEE	# # # # # # #	FF	ĒĒ	TIM TIM
41.88 26.85 17.81	25.92	19.26	16.07 0.00 0.00	25.83 11.92 10.41 0.40	26.06 18.62 10.68	9.53	0.00	36.88	24.37	16.09 29.76 0.11	10.46	0.02	21.90 8.26
9.19 × 6.05 × 0.00 ×	17.94	3.41 ×	3.03 0.08 0.08	6.65 1.69 0.00 0.00	8.82 × 5.72 × 1.88 ×	1.55 > 0.00	1.89	5.51 >	10.50 × 5.07 10.23 ×	4.66 0.00	3.77 >	0.00 >	4.10 >
3.56 > 0.00	0.0	0.0	888	0.0000	0.00	0.00	0.00	3.47	3.59 > 3.49	0000	0.00	0.00	0.00
320.00 × 320.00 ×	320.00	317.00 943.00	313.00 320.00 1000.00	320.00 320.00 1000.00	306.00 309.00 320.00 ×	30.90	309.00	309.00	320.00 × 1000.00 ×	320.00 1000.00 1000.00	3.09	9.40	320.00
A A A	A A	8.6	888	8888	300.00	0.00	0.0	89.00 0.00	89.00 × 286.00 × 284.00 ×	888	0.00	0.00	8.0
	3.46	0.00	0.00	0.00	14.60 7.75 1.88	0.00	0.00	7.47	10.50 7.07 10.23	5.03 19.98 0.00	3.77	6.78	0.00
320.00 × 320.00 × 1000.00	1000.00 >	315.00	159.00 93.50 279.00	320.00 > 320.00 > 1000.00	179.00 210.00 320.00 v	21.00	210.00	210.00	320.00 × 740.00 1000.00 ×	320.00 × 763.00 279.00	3.20 >	4.04	320.00 >
		0.00	0.00	48.10 v 190.00 v 0.00 v	12.30 27.10 170.00 >	0.00	0.00	28.10	30.60 × 105.00 97.80 ×	63.60 × 38.20 0.00	0.00	0.00	0.00
37.40 9.35 11.14	3.73	1.89	7.09 0.67 0.00	13.40 3.63 5.37 0.00	20.70 16.81 2.58	2.88	3.86	32.70	32.00 9.15 17.88	6.57 20.03 0.00	6.73	0.00	0.00
320.00 > 320.00 > 1000.0 >	97.70	140.00	73.10 89.80	320.00 × 320.00 × 791.00 × 561.60	108.00 155.00 320.00 >	15.50	155.00	155.00 108.00	320.00 × 530.00 1000.0 ×	296.00 394.00 151.00	1.55	1.73	320.00 >
8.55 34.20 >	13.50	8.8 8.7	0.00 0.00 0.00	23.90 × 88.10 × 147.00 0.00	5.24 9.22 124.00 >	5.38	40.20	4.75 54.20	10.00 × 57.90 × 55.90 ×	45.10 19.70 0.00	0.00	0.00	22.80 \$
0.886	0.643	0.935	1.023 0.914 0.952	1.023 0.850 0.952 0.551	0.817 0.576 1.087	0.588	0.606	0.554	0.554 0.771 0.592	0.362 0.588 0.526	0.751	0.814	0.469
	2 Z		SSEU	SNK	0 01H 82R 2XY	R6A	000 C	0 00M	X82	O CALL	SNZ C	SOO	90S 0
03/01/8 11/28/8 12/14/8	12/20/8	11/28/6	03/01/8 11/30/8 11/30/8	03/01/8 11/30/8 11/30/8 04/26/9	07/03/8 08/08/8 11/01/9	06/27/8 08/15/8	07/11/8	07/11/8 08/16/8	07/11/8 08/16/8 04/26/9	07/25/8 10/04/8 04/26/9	07/26/8	07/26/8	07/26/89 axs 11/30/89 so6
8 8 8	3 3	8 8	9 9 9	3 3 3 3	888	22	57	57 57	57 89	58 88	8 88	58 88	% %
5067 5067 5067	5067	5069 5069	507 507 508 508	8 8 8 8 8 8 8 8 8 8 8 8 8 8 8 8 8 8 8	5121 5121 5121	5128 5128	5136 5136	5137 5137	5138 5138 5138	518 518 518 518	5189 5189	5191 5191	5201 5201
	48 03/01/89 QUK 0.886 8.55 > 320.00 > 37.40 34.80 > 320.00 > 9.19 90.00 > 320.00 > 3.56 > 9.19 > 41.88 48 11/28/89 SWQ 0.535 34.20 > 320.00 > 9.35 79.10 > 320.00 > 4.05 280.00 > 320.00 > 1.14 > 4.05 > 26.85 48 12/14/89 SYF 0.723 89.70 > 1000.0 > 11.14 0.00 > 1000.00 0.00 > 1000.00 0.00 > 17.81	48 03/01/89 QUK 0.886 8.55 > 320.00 > 37.40 34.80 > 320.00 > 9.19 90.00 > 320.00 > 3.56 > 9.19 > 41.88 48 11/28/89 SWQ 0.535 34.20 > 320.00 > 9.35 79.10 > 320.00 > 4.05 280.00 > 320.00 > 1.14 > 4.05 > 26.85 48 12/14/89 SWF 0.723 89.70 > 1000.0 > 11.14 0.00 > 1000.00 0.00 0.00 0.00 0.00 17.81 48 12/20/89 TOG 0.643 13.50 640.00 47.40 35.70 > 1000.00 28.04 0.00 > 1000.00 0.00 17.94 25.92 48 03/01/89 QUL 0.828 26.10 97.70 3.75 52.30 181.00 3.46 0.00 > 320.00 3.20.00 1.87 17.90	48 03/01/89 QUK 0.886 8.55 > 320.00 > 37.40 34.80 > 320.00 > 9.19 90.00 > 320.00 > 3.56 > 9.19 > 41.88 48 11/28/89 SWQ 0.535 34.20 > 320.00 > 9.35 79.10 > 320.00 > 4.05 280.00 > 320.00 > 1.14 > 4.05 > 26.85 48 12/14/89 SVF 0.723 89.70 > 1000.00 > 11.14 0.00 > 1000.00 0.00 0.00 > 1000.00 0.00	48 03/01/89 CUK 0.886 8.55 > 320.00 > 37.40 34.80 > 320.00 > 9.19 90.00 > 320.00 > 3.56 > 9.19 > 41.88 48 11/28/89 SM 0.535 34.20 > 320.00 > 11.4 > 4.05 280.00 > 320.00 > 1.14 > 4.05 26.85 26.85 48 11/28/89 SM 0.535 34.20 > 320.00 > 11.14	48 11/28/99 SNP 0.535 34.20 > 37.40 34.80 > 320.00 > 9.19 90.00 > 320.00 > 3.56 > 9.19 > 41.88 4.05 ≥ 26.85 2.05 ≥ 26.85 4.05 ≥ 26.85 2.05 ≥ 26.85	48 03/01/89 aux 0.856 8.55 > 320.00 > 9.35 79.10 > 320.00 > 4.05 280.00 > 320.00 > 320.00 > 1.14 > 4.05 > 26.85 48 11/22/89 stq 0.535 34.20 > 320.00 > 9.35 79.10 > 320.00 > 4.05 280.00 > 320.00 > 11.4 > 4.05 > 26.85 48 12/12/89 stq 0.535 34.20 > 320.00 > 9.35 79.10 > 320.00 > 26.00 0.00 > 100.00 1.14 > 4.05 > 26.85 48 12/12/89 stq 0.535 34.20 > 9.10 > 11.14 0.00 > 100.00 0.00 > 100.00 0.00 > 11.40 0.00 > 17.50 48 11/12/89 stq 0.535 34.00 0.00 34.00 0.00 34.00 0.00 1.70 48 11/30/89 stq 0.535 10.30 75.2 41.20 20.00 31.50 0.00 1.70 48 11/30/89 stq 0.535 10.30 75.10 7.09 22.60 150.00 0.00 31.50 0.00 1.70 48 11/30/89 stq 0.535 10.30 75.10 1.89 0.00 21.50 0.00 0.00 0.00 0.00 0.00 0.00 0.00 0.00 0.0	48 11/20/89 SMG 0.586 8.55 > 320.00 > 37.40 34.80 > 320.00 > 9.19 90.00 > 320.00 > 3.56 > 9.19 > 41.88 4.85 > 11/20/89 SMG 0.533 34.50 > 9.15 > 9.10 > 320.00 > 9.15 9.00 > 320.00 > 3.50 > 9.19 > 41.88 4.85 > 9.19 > 4.05 > 280.00 > 320.00 > 1.14 > 4.05 > 28.65 9.19 > 4.10 > 100.00 9.00 9.10 9.10 9.10 9.00 9.10 9.	48 12/20/89 CMC 0.886 8.55 > 320.00 > 37.40 34.80 > 320.00 > 9.19 90.00 > 320.00 > 1.45 > 4.05 \$20.00 > 1.45 > 4.188	48 11/20/99 940 0.535 34.00 > 37.40 35.40 > 37.40 94.00 > 5.19 90.00 > 320.00 > 3.56 > 9.19 > 41.48	4.8 11/20/99 SMC 0.585 34.20 > 372.00 > 37.40 34.80 > 320.00 > 4.05 280.00 > 320.00 > 114.5 4.05 > 26.85 4.89 4.8 11/20/99 SMC 0.5723 89.70 > 100.00 > 17.44 0.00 > 100.00 > 100.00 > 114.5 4.05 > 26.85 4.8 11/20/99 SMC 0.5723 89.70 100.00 > 17.54 100.00 100.00 > 10.00 > 114.5 4.05 > 26.85 4.8 11/20/99 SMC 0.5723 18.70 100.00 1.57 100.00 1.50 100.00 10.00 11.50 100.00 1.50 100.00 1.50 100.00 1.50 100.00 1.50 100.00 1.50 100.00 1.50 11.50 1.50	44 11/23/69 SHC 0.855 34.000 9.35 77.40 34.48 9.20.00 9.40 0.000 130.00 11.6 4.05 5.648 11.2 4	4.8 17/28/99 SMR 0.535	48 17/20/99 DMR 0.355 5.720 00 > 97.40 14.80 > 302.00 > 9.19 90.00 > 350.00 > 1.14 > 4.05 26.54 14.80 14.80 14.80 48 12/70/99 DMR 0.355 5.720 90 > 91.71 49 12/70/99 DMR 0.355 14.20 40 12/70/99 DMR 0.355 14.20

Table 18 (Cont'd)

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Assay	FF	FF	EE	ĒĒ	FE	EEE		FF	FF	FFF	EEE		ĔĒ	
I¥	3.8 3.8	29.23 19.06	42.33	5.95 18.23	3.56	0.00 0.90 1.91	19.39 0.00 5.60	10.97	0.00	10.51 11.78 5.24	3.54 15.99 17.43	36.83 24.98	5.04	22.03 18.86 15.69 18.32
15	0.00	5.20 >	.22.00 >	3.13	5.00	0.00	3.57 × 0.00 1.71	1.08 >	1.16 > 0.00	0.00	2.24 × 5.64 × 5.73 ×	8.12 > 4.48 > 2.83 >	1.22 > 0.00 >	3.71 × 2.23 × 1.96 × 6.81 ×
AI 95	9.0	0.00	0.00	0.00	9.35	00.00	0.00	0.00	0.0	9000	0.00	3.11	0.00	69.80 × 0.00 × 3.37
26 21	30.80	320.00 958.00	320.00 1000.00	320.00	276.00	6.11 3.20 7.38	320.00 100.00 43.30	320.00 320.00	92.40 87.90	320.00 320.00 10.00	100.00 100.00 8.00	1000.00 973.00 320.00 ×	320.00 1000.00	189.00 32.00 100.00 ×
36 31	9.0	° 0.0	0.00	0.00	29.50	888	0.00	0.00	9.0	0.00	0.00	0.00 × 313.00 288.00 ×	0.00	29.60 \$ 0.00
AI 50	2.23	5.20	22.00	0.00	4.38	2.00 0.00 0.00	6.14 0.00 2.64	3.34	0.00	0.00 0.00	5.34 34.67 8.33	8.12 6.04 2.83	1.22	6.42 2.23 4.79 9.17
1C 50	9.77	320.00 >	320.00 >	183.00	62.50	1.73 × 1.00 0.76	27.40 25.90 19.70	18.30	24.90	320.00 86.40 10.00	5.34 × 18.60 0.27 ×	1000.00 > 663.00 > 320.00 >	320.00 >	6.42 × 32.00 × 71.40 66.00
10 50	9.12	61.60 >	14.50 >	0.00	14.30	0.00	4.47	5.47 0.00	9.57	21.00	1.00	123.00 > 110.00 113.00 >	263.00 > 0.00	1.00 14.40 × 14.90 7.20
AI 25	3.45	9.53	61.40	1.44	3.10	0.00	8.12 0.00 2.61	4.52	2.01	11.80 4.03 0.00	2.24 < 9.45 5.73 <	42.99 11.22 7.61	9.1.	3.71 < 5.09 2.87 153.13
70 25	4.11	320.00 >	320.00 >	7.53	25.20	1.00 - 0.48	16.00 8.57 12.80	5.89	11.10	320.00 > 55.50 7.14	2.24 > 3.02 > 0.18 >	1000.0 > 492.00 320.00 >	320.00 >	3.71 × 32.00 × 29.10 × 49.00 ×
10 25	0.00	33.60 >	5.21 >	3.06	8.08	6.00	1.96	1.30	5.53	27.10 × 13.80 0.00	0.32	23.30 × 43.80 × 42.10 ×	161.00 > 247.00	1.00 6.29 × 10.20 0.32
Diff.	0.421	0.721	0.798	0.656	1.185	0.816 < 0.653 0.701	1.091 0.526 0.596	0.693	0.561	0.732	0.732 < 0.646 < 0.496 <	0.751	0.676	0.550 < 0.786 0.795 0.530 <
ž*	07/26/89 axv 11/30/89 so2	89 OZA 89 SYD	89 028 89 10G	89 P79 97 T70	89 PHT 90 TF5	89 P02 90 V2X 90 XAT	89 PHS 90 VMV	89 PKG 90 VMV	89 P27	89 P04 90 V2Y 90 XCY	89 P04 90 V22 90 XLB	89 SWA 89 SYB 90 VZZ	90 W01	89 PKR 89 SYC 90 M03
Test	07/26/ 11/30/	03/08/89	03/08/89 12/20/89	03/21/89	04/12/89	03/28/89 05/08/90 07/06/90	04/12/89 04/26/90 07/06/90	04/18/89 04/26/90	05/05/89 06/07/89	03/28/89 05/08/90 07/06/90	03/28/89 05/08/90 07/17/90	12/12/89 12/14/89 05/08/90	05/08/90 07/06/90	04/18/89 12/14/89 05/08/90 07/17/90
Ship-	5 50	25	22	22	22	288	288	28	22	288	288	223	33	2233
AVS.	5207 5207	5241	5242 5242	5283 5283	5350 5350	5405 5405 5405	5450 5450 5450	5452 5452	5459	5482 5482 5482	2483 2483 5483	222	5489	2222 2223 2233 2233

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EE EE	ĒĒ
8.62 5.18 60.74 1.06	0.0 0.0 0.0
2.14 0.79 0.00	0.00
0.00 0.00 0.00 0.00	13800 • 0.00
90.30 32.00 320.00 >	100,00
A A A	7.26 > 0.00 >
3.18 1.31 151.00 0.00	0.00
7.32 3.01 320.00 > 715.00	100.00
2.30 2.30 0.00 4	0.00
	100000 < 0.00
	100000 >
1.49	0.00
0.658 1.123 0.898 0.759	0.392 < 0.392
789 R4H 790 ZXZ 789 QEF 789 R4J	789 PYE 789 PYE
08/09, 11/01, 06/14, 08/09,	05/03/89 05/03/89
	22
5539 5539 5546 5546	*5580 5580
	56 08/09/89 R4H 0.658 1.49 4.92 3.30 2.30 7.32 56 11/01/90 2x2 1.123 0.71 1.81 2.53 2.30 3.01 56 06/14/89 QEF 0.898 1.39 > 320.00 > 230.00 2.11 > 320.00 56 08/09/89 R4J 0.759 0.00 518.00 0.00 0.00 715.00

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EEEE	FFF	ĔĔ	ĒĒ	ĒĒ	ĒĒ	ĒĒ	EE	T I	T E	F F	ĒĒ	ËĒ	ĔĔ	ĒĒ	
6.43 40.14 6.03 35.07	9.80 16.92 0.00	28.90	6.86	30.39	4.76	5.35	7.19	18.47	18.81 28.34	13.22	28.65	0.03	36.45	3.28	
1.86 14.77 > 0.00 > 10.14	3.61	5.66 0.00	0.00	3.94	0.69 •	9.0	1.79	3.65 >	2.41 > 7.69	2.27	37.65	2.00	0.00	0.00	
0.00	0.00	3.39 >	0.00	3.48	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	3.39	
1000.00 3200.00 > 1000.00 3200.00	320.00 320.00 920.00	320.00 >	320.00	318.00	320.00	320.00	320.00 882.00	320.00	320.00	30.70	320.00 320.00	9.55	9.67	32.00	
26.08 0.08 0.08 0.08	0.00	94.50 >	0.00	91.20	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	28.50	
3.40 16.32 0.00 15.85	5.32 0.00	5.66 0.00	0.00	5.38	1.48	9.00	2.56	3.85	2.41	3.40	50.34	0.00	0.00	0.00	
1000.00 × 3200.00 × 1000.00 × 2630.00	320.00 280.05 167.00	320.00 >	320.00	215.00	110.00	186.00	251.00	320.00 > 813.00	320.00 > 758.00	19.20	320.00 >	5.47	5.94	21.60	
294.00 v 196.00 v 0.00 v 166.00	0.00 × 52.70 0.00	56.60 \$	0.00	39.98 8.73	74.90	0.00	97.80	83.10 > 74.80	133.00 > 61.10	5.66 0.00	6.36 > 8.07 >	2.68	0.00	10.00	
36.98	4.63 9.67 0.00	7.52	2.73	24.23	1.06	1.92	3.13	7.47	5.76	4.36	53.06	3.77	0.00	11.62	
547.00 2900.0 715.00 1680.0	320.00 > 190.00 8.30	320.00 >	192.00	157.00 157.00	52.00 150.00	90.60	175.00	320.00 >	320.00 >	12.80	239.00	3.20	3.86	49.00	
78.30 217.00 114.00	69.10 × 19.70 0.00	42.50 >	0.00	67.9	49.00	47.20	55.90	42.80 >	55.60 ×	2.94	4.51	0.00	0.00	4.22	
0.835 0.744 0.613 0.907	0.739	0.548	0.976	0.785	0.714	0.785	0.776	0.798	0.722	0.624	0.681	0.815	0.706	0.715	
99 SSU 90 TVG 90 WHO 90 ZYO	PR RBG	89 QTQ 89 R8A	789 SMT	89 SMU 90 TGP	89 SN9	89 SNB	789 SOC 790 TGP	'89 SE2	789 SE9	89 SAS	89 SAU 790 TLN	39 SOK 790 TGU	89 SON 190 TGX	39 SOX	
12/05/ 4 02/08/ 05/31/ 7 11/01/	07/06/ 08/16/ 05/01/	07/19, 08/16,	11/29,	11/29,	11/29/	11/29,	11/30,	11/03,	11/03/	11/29,	11/29,	11/30,	11/30,	11/30,	
62 GABSN 67 62/67	57 57 65	57	59	20	20	20	33	33	33	22	22	33	33	33	
\$601 \$601 \$601	5643	5691 5691	5765 5765	5780 5780	5818 5818	5823 5823	5842	\$855 \$855	5869	5905	5910 5910	5917 5917	2924	2940	
	62 12/05/89 \$SU 0.835 145.00 547.00 3.78 294.00 > 1000.00 > 3.40 0.00 > 1000.00 0.00 1.86 6.43 6.43 6.70 1 0.00 > 10.00 > 10.00 > 10.00 > 10.00 > 10.00 > 10.00 > 10.00 > 14.77 > 40.14 6.77 > 40.14 6.77 > 40.14 6.77 > 40.14 6.77 > 40.14 6.77 > 40.14 6.77 > 40.14 6.77 > 40.14 6.77 > 40.14 6.77 > 40.14 6.77 > 40.14 6.77 > 40.14 6.00 > 10.00 0.00 0.00 0.00 0.00 0.00 0.0	62 12/05/89 \$\$U 0.835 145.00 547.00 3.78 294.00 > 1000.00 > 3.40 0.00 > 1000.00 0.00 1.86 6.43 6.43 6.208 0.744 78.30 2900.0 34.96 196.00 > 3200.00 > 16.32 846.00 > 3200.00 > 3.78 14.77 > 40.14 6.70 5.71 217.00 715.00 3.29 0.00 > 1000.00 0.00 0.00 0.00 0.00 0.00	42 12/05/89 \$\$\text{6.02}\$ 12/05/89 \$\$\$\text{6.02}\$ 12/05/89 \$\$\$\text{6.02}\$ 12/05/89 \$\$\$\text{6.02}\$ 12/05/89 \$\$\$\text{6.02}\$ 12/05/89 \$\$\$\text{6.02}\$ 2900.0 \$\$\text{6.02}\$ 3.78 \$\$\text{6.00}\$ 290.00 \$\$\text{6.02}\$ 3.50 \$\$\text{6.00}\$ 3.200.00 \$\$\text{6.03}\$ 3.78 \$\$\text{6.017}\$ 40.14 \$\$\text{6.07}\$ 40.14 \$\$\text{6.07}\$ 40.14 \$\$\text{6.02}\$ 25/00.00 \$\$\text{6.00}\$ 0.00 \$\$6	62 12/05/89 SSU 0.835 145.00 547.00 3.78 294.00 > 1000.00 > 3.40 0.00 > 1000.00 0.00 1.86 6.43 6.43 6.400 > 3.78 14.77 > 40.14 6.83 02/08/90 TVG 0.744 78.30 2900.0 34.96 196.00 > 3200.00 > 16.32 846.00 > 3200.00 > 3.78 14.77 > 40.14 6.70 5715.00 7715.00 7715.00 3.29 0.00 > 1000.00 0.00 0.00 0.00 0.00 0.00	62 12/05/89 SSU 0.835 145.00 547.00 3.78 254.00 > 1000.00 16.32 846.00 > 3200.00 3.40 0.000 > 1000.00 1.85 64.37 40.14 CABSN 02/08/90 TVG 0.744 78.30 2900.0 36.96 196.00 > 1000.00 16.32 846.00 > 3200.00 3.78 14.77 40.14 67 05/31/90 MM0 0.613 217.00 715.00 3.29 0.00 1000.00 0.00 1000.00 0.00 10.00 0.	62 12/05/89 Ssu 0.835 145.00 547.00 3.78 294.00 > 1000.00 1.6.32 646.00 > 1000.00 0.00 1.685 64.33 GABSN 02/08/90 TvG 0.744 772.00 34.96 196.00 > 1000.00 16.32 646.00 > 1000.00 3.78 14.77 40.14 67 05/31/90 WHO 0.613 217.00 715.00 3.29 0.00 > 1000.00 16.32 646.00 > 1000.00 0.00 1.77 40.14 67/67 71/10/190 ZvG 0.907 11/20/190 WHO 0.613 11/20 14.70 166.00 2630.00 15.85 0.00 2000.00 10.00 0.00 10.00 0.00 10.00 0.00 10.00 0.00 10.00 0.00 10.00 0.00 10.00 0.00 <th< th=""><th>62 12/05/99 SSU 0.633 145.00 547.00 3.78 294.00 > 1000.000 3.40 0.00 > 1000 0.00 1.86 6.43 6488 02/88/90 TVG 0.74 78.30 2990.0 35.99 196.00 > 1000 16.32 66.00 > 3200.00 3.78 14.77 40.14 67 0.513/90 WMO 0.613 217.00 715.00 18.29 10.00 <th< th=""><th>42 12/05/99 SSU 6.835 145.00 547.00 3.78 294.00 > 1000.00 3.40 0.00 > 1000.00 1.86 6.43 46.88 02/08/90 TWG 0.744 774 774 77.00 1.77 4.14 6.06 11/2/190 TWG 0.613 11/2 12.00 35.00 12.30 10.00 10.00 10.00 1.77 4.14 62/67 11/11/190 ZWG 0.507 11/2 14.00 14.00 16.00 16.00 10.00 1</th><th>42 12/05/99 58.0 0.835 145.00 547.00 3.78 294.00 3.40 0.00 1000.00 1.00 1000.00 1.00</th><th>ACT ACT ACT</th></th<><th>42 12/05/99 \$530 143.00 54.70 3.78 145.00 520.00 3.40 0.00 1000 1000 1000 1.00<th>42 12/05/96 stall 6.35 445.00 477.00 3.78 294.00 1000.00 3.40 1.186 6.41 464.00 1000.00 1.50 1.186 0.00 1000.00 1.00 1.00 4.01 62/47 11/10/100 trol 6.31 17.50 175.00 18.20 14.70 16.00 10.0</th><th>42.04 14704/99 stal 14320 45.0 45.0 35.0</th><th>44. Control (1) C</th><th> 1.5 1.5</th></th></th></th<>	62 12/05/99 SSU 0.633 145.00 547.00 3.78 294.00 > 1000.000 3.40 0.00 > 1000 0.00 1.86 6.43 6488 02/88/90 TVG 0.74 78.30 2990.0 35.99 196.00 > 1000 16.32 66.00 > 3200.00 3.78 14.77 40.14 67 0.513/90 WMO 0.613 217.00 715.00 18.29 10.00 <th< th=""><th>42 12/05/99 SSU 6.835 145.00 547.00 3.78 294.00 > 1000.00 3.40 0.00 > 1000.00 1.86 6.43 46.88 02/08/90 TWG 0.744 774 774 77.00 1.77 4.14 6.06 11/2/190 TWG 0.613 11/2 12.00 35.00 12.30 10.00 10.00 10.00 1.77 4.14 62/67 11/11/190 ZWG 0.507 11/2 14.00 14.00 16.00 16.00 10.00 1</th><th>42 12/05/99 58.0 0.835 145.00 547.00 3.78 294.00 3.40 0.00 1000.00 1.00 1000.00 1.00</th><th>ACT ACT ACT</th></th<> <th>42 12/05/99 \$530 143.00 54.70 3.78 145.00 520.00 3.40 0.00 1000 1000 1000 1.00<th>42 12/05/96 stall 6.35 445.00 477.00 3.78 294.00 1000.00 3.40 1.186 6.41 464.00 1000.00 1.50 1.186 0.00 1000.00 1.00 1.00 4.01 62/47 11/10/100 trol 6.31 17.50 175.00 18.20 14.70 16.00 10.0</th><th>42.04 14704/99 stal 14320 45.0 45.0 35.0</th><th>44. Control (1) C</th><th> 1.5 1.5</th></th>	42 12/05/99 SSU 6.835 145.00 547.00 3.78 294.00 > 1000.00 3.40 0.00 > 1000.00 1.86 6.43 46.88 02/08/90 TWG 0.744 774 774 77.00 1.77 4.14 6.06 11/2/190 TWG 0.613 11/2 12.00 35.00 12.30 10.00 10.00 10.00 1.77 4.14 62/67 11/11/190 ZWG 0.507 11/2 14.00 14.00 16.00 16.00 10.00 1	42 12/05/99 58.0 0.835 145.00 547.00 3.78 294.00 3.40 0.00 1000.00 1.00 1000.00 1.00	ACT ACT	42 12/05/99 \$530 143.00 54.70 3.78 145.00 520.00 3.40 0.00 1000 1000 1000 1.00 <th>42 12/05/96 stall 6.35 445.00 477.00 3.78 294.00 1000.00 3.40 1.186 6.41 464.00 1000.00 1.50 1.186 0.00 1000.00 1.00 1.00 4.01 62/47 11/10/100 trol 6.31 17.50 175.00 18.20 14.70 16.00 10.0</th> <th>42.04 14704/99 stal 14320 45.0 45.0 35.0</th> <th>44. Control (1) C</th> <th> 1.5 1.5</th>	42 12/05/96 stall 6.35 445.00 477.00 3.78 294.00 1000.00 3.40 1.186 6.41 464.00 1000.00 1.50 1.186 0.00 1000.00 1.00 1.00 4.01 62/47 11/10/100 trol 6.31 17.50 175.00 18.20 14.70 16.00 10.0	42.04 14704/99 stal 14320 45.0 45.0 35.0	44. Control (1) C	1.5 1.5

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			ĦĦ	E E	EE	FFF	FE	ĔĔ	ĒĒ	ĒĒ	EE	ĒĒ	ĒĒ	EE	ĒĒ	
0.02	70.07	0.00 29.64 29.21	10.47	3.00	11.64	51.70 22.19 15.81	17.11	22.36	19.21	45.62	27.69	26.05	44.15	53.82	21.76	
0.00	0.00	9.75 7.13 v	0.00	0.00 >	1.13 > 0.00	19.85 × 3.09 × 2.92	0.06	1.73	0.00 \$	8.69 > 2.46	4.15 > 2.76 >	4.09 > 2.21	17.51 > 12.84 >	21.52 > 2.27 >	3.00 >	
8.8	3.0.5	0.00	9.0	0.0	0.00	1.90 × 3.24 3.22	1.07	1.06 v	0.00	3.54 >	3.40	1.15 > 3.07	10.63	2.22 > 1.09 >	1.12 > 3.39	
	320.00	320.00 32.00 ×	320.00	303.00	320.00	320.00 > 966.00 966.00	320.00 > 935.00	320.00 >	320.00 320.00	320.00 > 963.00	98.60 97.80	320.00 >	320.00 >	320.00 >	320.00 >	
8.6	A A		0.00	0.00	0.00	A	A	A A	0.00	A	30.00	A	30.10 > 70.40 >	A A	A A	
0.4	0.00	17.99	0.00	0.00	1.13	19.85 4.16 3.94	2.05	£::	0.00	3.50	5.62 3.84	3.31	18.44	21.52	3.00	
13.40	20.50	30.90	303.00	153.00	320.00 >	320.00 × 660.00 660.00	320.00 > 346.00	320.00 >	320.00	320.00 > 626.00	67.10 64.90	320.00 >	320.00 >	320.00 >	320.00 > 938.00	
0.00	0.00	0.00	0.00	0.00	282.00 > 0.00	16.10 × 159.00	156.00 > 313.00	179.00 > 288.00 >	0.00 >	36.80 >	11.90	78.20 >	17.30 >	14.90 >	107.00 > 141.00	
1.10	0.00	0.00 13.07 9.81	3.65	1.09	2.49	73.07	0.09	42.55	25.64	3.29	3.80	2.8%	23.77	3.8	5.45	
9.60	7.12	2.38 17.30 13.30	158.00	73.40	320.00 > 635.00	320.00 × 490.00 490.00	10.00	320.00 >	320.00 >	320.00 >	46.50	320.00 >	304.00	320.00 >	320.00 > 629.00	
6.01	0.00	1.33	0.00	67.50	129.00 >	4.38 > 112.00 121.00	109.00	7.52 >	12.50 >	12.90 >	5.09	36.20 >	12.80	4.21 > 82.00 >	58.70 × 85.50	
6773	0.636	0.636 0.523 0.823	0.675	0.731	0.668	0.851 1.007 0.870	1.007	0.801	0.763	0.763	0.755	1.019	0.875	0.931	0.931	
	89 SF1	89 SF1 90 TNO 7V7	89 SFM 90 TNO	SP SHK	39 SHO 90 TNQ	20 ZVR 20 ZVR 20 103	20 TAB 20 ZVR	00 TAD 00 103	00 TAE	90 TAE	20 TAF 20 2VT	00 TAH 00 ZVU	90 TC6	90 TC7 90 ZY2	90 TC7 90 196	
12/01/8	11/07/1	11/07/102/02/01/02/08/9	11/07/	11/09/1	11/09/1	01/11/ 10/30/ 12/05/	01/11/5	01/11/	01/11/	01/11/	01/11/	01/11/	01/16/	01/16/	01/16/	
2.2	2 2 2	222	25	22	22	622	2 29	2 2	62	29	62 62	62	82	62	62 62	
	288	5998 5998 5998	6011	6209	6041	6195 6195 6195	6196	9500	6201	6202	6204	6207	6218 6218	6219	6220 6220	
	61 12/01/89 SDT 0.473 6.01 6.60 1.10 0.00 13.40 0.00 0.00 30.50 0.00 0.00 0.02	61 12/01/89 SDT 0.473 6.01 6.60 1.10 0.00 13.40 0.00 0.00 30.50 0.00 0.00 0.02 61 02/01/90 TNW 0.563 5.48 8.21 1.50 9.38 16.30 1.74 0.00 30.40 0.00 0.88 1.97 61 11/07/89 SF1 0.636 0.00 7.12 0.00 0.00 20.50 0.00 0.00 320.00 0.00 0.00 0.07 51 11/01/90 ZV1 0.861 1.31 7.79 5.03 1.77 25.10 14.17 3.02 > 100.00 > 33.46 4.41 > 23.56	61 12/01/89 SDT 0.473 6.01 6.60 1.10 0.00 13.40 0.00 0.00 30.50 0.00 0.00 0.00 0.02 61 02/01/90 TNW 0.563 5.48 8.21 1.50 9.38 16.30 1.74 0.00 30.00 0.00 0.00 0.08 1.97 61 11/07/89 SF1 0.636 0.00 7.12 0.00 0.00 20.50 0.00 0.00 320.00 0.00 0.00 0.00 0.00 61 11/07/89 SF1 0.636 0.00 2.38 0.00 0.00 7.23 0.00 0.00 320.00 0.00 0.00 0.00 0.00	61 12/01/89 SDT 0.473 6.01 6.60 1.10 0.00 13.40 0.00 0.00 30.50 0.00 0.00 0.00 1.97 (1.77 0.00) 1.74 0.00 30.40 0.00 0.00 0.08 1.97 (1.77 0.00) 1.74 0.00 30.40 0.00 0.00 0.00 0.00 0.00 0.	61 12/01/89 SDT 0.473 6.01 6.60 1.10 0.00 13.40 0.00 0.00 30.50 0.00 0.00 0.08 1.97 (61 12/01/89 SPT 0.563 5.48 8.21 1.50 9.38 16.30 1.74 0.00 30.00 30.40 0.00 0.00 0.00 0.00 0	61 12/01/89 SDT 0.473 6.01 6.60 1.10 0.00 13.40 0.00 0.00 30.50 0.00 0.00 0.00 0.00 0.	61 12/01/89 SDT 0.473 6.01 6.60 1.10 0.00 13.40 0.00 30.50 0.00 0.00 0.00 0.00 1.97 1.97 1.00 0.00 0.00 0.00 0.00 0.00 0.00 0.0	61 12/01/89 SDT 0.473 6.01 6.66 1.10 0.00 13.40 0.00 0.00 30.50 0.00 0.00 0.00 1.74 0.00 30.40 0.00 0.00 0.00 1.97 1.97 0.00 30.40 0.00 0.00 0.00 0.00 1.74 0.00 30.40 0.00 0.00 0.00 1.97 1.97 0.00 1.174 0.00 30.40 0.00 0.00 0.00 0.00 1.97 0.00 0.00 0.00 0.00 0.00 0.00 0.00 0	61 11/07/89 SPI 0.435 6.01 6.60 1.10 0.00 13.40 0.00 0.00 30.50 0.00 0.00 0.00 1.97 11/07/89 SPI 0.635 5.48 6.21 1.50 9.38 16.30 1.77 0.00 0.00 0.00 320.00 0.00 0.00 0.00 1.10 0.00 11/01/90 ZVI 0.861 1.31 7.79 5.93 1.77 25.10 14.17 3.02 100.00 33.16 4.41 > 23.56 11/01/90 ZVI 0.861 1.31 7.79 5.93 1.77 25.10 14.17 3.02 > 10.00 > 320.00 0.00 0.00 0.00 0.00 0.00 0.00 0.	61 11/07/99 \$FI 0.635	64 12/01/99 STI 0.473 6.01 6.40 1.10 0.00 13.40 0.00 0.00 30.50 0.00 0.	64 11/07/99 ST	1,007/89 51 0.473 6.01 6.60 1.10 0.00 13.40 0.00 0.00 30.50 0.00 0.	11/07/99 51 12/07/99 51 12/07/99	11/07/99 ST 0.457 0.40 0.00	11/07/99 St. 0.553 C. 10 C. 10

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Assay Type	EE	EEEE	EEE	ĒĒ	EE	EE	EE	EE	EE	ĔĒ	ĒĒ	ĒĒ	ĒĒ	ĒĒ	ĔĒ	
TAI	16.91 15.38	1.45 11.69 4.19 0.60	0.76 12.02 3.45	3.80	14.05	1.83	4.07	6.31	4.55	16.63	3.30	11.96	6.69	9.98	3.56	
ıs	1.92 > 2.74 >	0.000	1.04 >	1.14 > 0.00 >	2.55	1.42	0.41	0.00 > 2.22 >	1.38	0.00	0.00	1.47 > 0.00	4.95	8.81	5.95 >	
A1 95	1.07 >	0.000	0.00	0.0	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	3.50	0.0	0.00	
70 95	320.00 >	1000.00 3200.00 3200.00 3200.00	1000.00 3200.00 3200.00	1000.00 3200.00	320.00 951.00	30.80	309.00	320.00	163.00	320.00 1000.00	320.00 320.00	320.00 966.00	320.00	96.60 268.00	320.00 1000.00	
S6 21	300.00 >	^ ^ ^ ^ 00000	0.00	0.00	0.00	0.00	0.00	0.00 × 2950.0 ×	0.00	0.00	0.00	0.00	0.00 >	0.00	0.00	
AI 50	1.92	0.00	1.04	0.00	3.71	0.00	2.34	0.00	3.42	0.81	0.00	1.47	0.00	0.00	0.00	
TC 50	320.00 >	1000.00 3200.00 3200.00 3200.00	1000.00 3200.00 > 3200.00 >	1000.00 > 3200.00	269.00	19.50	182.00	320.00	59.40	320.00 >	320.00	320.00 >	320.00	66.00	320.00 882.00	
10 50	166.00 \$	1600.00 v 0.00 v 0.00 v	0.00 > 3080.00 > 2690.00 >	840.00 × 0.00 ×	0.00	9.30	7.60	0.00 >	25.90	395.00 >	0.00 > 259.00 >	217.00 > 0.30	0.00 >	0.00	9.73	
AI 25	3.66	1.61 6.34 2.50 0.00	1.13 2.65 2.04	1.84	5.29	2.61	0.00	0.00	2.34	11.67	2.28	2.70	3.62	1.72	0.00	
35 JI	320.00 >	3200.0 > 1440.0 > 3200.0	3200.0 > 3200.0 >	955.00	185.00 283.00	13.30	32.00	320.00	35.80	320.00 >	198.00	320.00 >	320.00 > 507.00	49.00	239.00	
10 25	120.00 >	621.00 × 505.00 × 576.00 × 0.00 ×	885.00 > 1210.0 > 1570.0 >	519.00	34.90	5.07	45.80	0.00 >	15.30	27.40 >	86.70 161.00 >	118.00 \$	88.50 × 51.30	3.28	5.18	
Diff.	0.973	0.836 0.814 0.656 0.900	0.733	0.803	0.925	0.464	0.956	0.956	0.970	0.548	0.740	0.724	0.592	0.592	0.690	
Test Pit Date #	01/16/90 TCA 11/01/90 2Y2	12/05/89 SSV 02/08/90 TVH 05/31/90 WHP 11/01/90 ZY3	12/05/89 SSW 02/08/90 TVI 05/31/90 WHQ	11/14/89 SJ8 02/06/90 TRP	11/14/89 SJB 02/06/90 TRR	02/15/90 U24 03/15/90 UNB	02/20/90 U32 03/20/90 UPC	02/20/90 U3Z 03/15/90 UNE	02/20/90 U40 03/20/90 UPD	02/22/90 USV 03/20/90 UPE	02/27/90 U7R 03/20/90 UPH	02/27/90 U71 03/20/90 UPI	03/01/90 U94 03/22/90 URG	05/10/90 W2K 07/06/90 XD1	05/10/90 WZN 07/06/90 XD2	
Ship- ment	33	62 1 6ABSN 0 67 0 67 1	62 1 GABSN 0 67 0	-0	10	55	55	63 63	55	22	25	22	63	33	33	
AVS SI	6225	6234 6234 6234 6234	6236 6236 6236	6243	6576	6308	63%	6335 (6337	6362	6374 6	6379	6391	6412	417	

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Assay Type	ËË	ĒĒ	ĒĒ	ĒĒ	ĒĒ	ĒĒ	FF	ĔĒ	EE	ĒĒ	EE	EE	FF	FIN	ĒĒ	EE	
IĀI	9.55	3.80	5.27	15.38	10.57	24.65	17.14 0.33	9.14	21.09	22.06	1.22	20.0%	24.26 18.05	1.97	4.82	25.97	
75	3.48	0.00	1.88 \$	1.70	2.72 > 0.00	4.0%	2.17 >	1.21 \$	0.00 >	4.56 > 2.07 >	2.18	4.72 >	3.03 >	0.00 >	0.00	4.62 0.00	
A1 %	0.00	0.00	0.00	0.00	0.00	1.15 > 0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	3.41	
26 21	320.00	90.30	88.20 10.00	320.00 1000.00	320.00	320.00 >	%.% %.%	320.00 1000.00	299.00	163.00	95.80 97.30	94.00 30.80	%.% %.%	320.00 320.00	320.00	320.00 >	
اد % اد %	0.00	0.00	0.00	0.00	0.00	278.00 > 0.00 >	0.00	0.00 \$	0.00	0.00	0.00	0.00	0.00	0.00 \$	0.00	93.90 >	
A1 50	0.00	0.00	9.51	2.39	4.15	70.0	2.92	1.21	19.91	7.01	0.00	8.35	3.04	0.00	0.00	67.9	
TC 50	247.00	26.90 63.8 0	9.51 ×	239.00	320.00 > 713.00	320.00 > 601.00	66.00 50.00	320.00 > 633.00	112.00	62.50 65.10	57.50 66.30	40.40	66.00 66.00	320.00 320.00 >	320.00	236.00	
10 50	0.00	0.00	0.00	100.00	7.10 \$	79.20 v 0.00	22.60	264.00 >	0.00	8.92	0.00	0.00	16.20 21.70	0.00 > 250.00 >	0.00 \$888.00 \$	36.30	
AI 25	3.68	3.55	1.88 < 2.62	7.65	6.15	7.72	8.53 0.00	2.43	17.62 23.94	40.72 22.15	0.00	9.71	20.13	0.00	0.00	13.18	
1C 2S	170.00	18.50	1.88	170.00	210.00	320.00 > 257.00	49.00	320.00 >	68.20 97.40	40.70 >	36.30	22.80	49.00	320.00	0.96	168.00	
10 25	46.20	12.90	1.00	22.20	34.20	41.50 >	5.74	132.00 >	3.87	1.00	0.00	2.35	5.33	0.00 >	201.00	12.70	
Diff.	0.701	0.688	0.527 < 1.066	0.583	0.583	0.504	0.504	0.587	0.728	0.732 < 0.588	0.731	0.549	0.567	0.733	0.790	0.333	
Test Pit	MON 06/90/90 05/06/90 NON	06/06/90 WP3 07/06/90 XD5	03/01/90 U99 03/22/90 URH	03/01/90 U9A 03/22/90 URH	03/01/90 U9A 03/22/90 URI	03/01/90 U9B 03/22/90 URI	03/01/90 U9B 03/22/90 URJ	03/01/90 U9D 03/22/90 URJ	06/06/90 WP7 07/10/90 XF8	06/06/90 WP9 07/10/90 XF9	05/17/90 W7T 07/10/90 XF9	05/17/90 W81 07/10/90 XF1	05/17/90 W83 07/10/90 XF3	05/23/90 W92 07/10/90 XF4	05/24/90 WCB 07/10/90 XF6	03/15/90 UNG 05/03/90 VAV	
Ship- ment	33	33	33	23	63	63	22	63	33	33	33	33	33	33	33	33	
AVS Ho.	6422	6436	0779	2445	6443	777	6445	6773	22	477	6482	7679	6501	6521	6560	6580	

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Assay Type	FF	E E	E E	ĒĒ	ĔĒ	ĔĔ	H	E E	ĒĒ	ËË	ĒĒ	H H	FFF	ĒĒ	FIE	H H
¥	10.54	9.27	12.31	29.32	6.28	21.14	7.77	0.00	28.40	13.42	5.43	3.31	3.78 20.67 3.83	16.56	6.87 5.88 0.90	14.37
2	0.00	1.59 >	2.59 >	11.03 >	1.8	3.39 >	1.57 >	0.39 >	3.02 >	0.87 > 2.85	3.40	1.12	0.71 v 4.22 0.00 v	3.92	0.00	3.43 > 1.85 >
A1 %	0.00	° 0.0 0.00	0.0	0.00	0.0	0.00	0.00	0.00	0.0	0.00	0.0	0.0	0.00	0.0	0.00	0.00
70 %	320.00	1000.00	1000.00	1000.00	30.90	320.00	320.00	320.00	309.00	320.00 >	320.00	9.91	320.00 1000.00 320.00	320.00 310.00	320.00 1000.00 320.00	320.00
35 25	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.0	289.00 >	0.00	0.00	0.00	0.00	0.00	0.00
A1 50	9.0	1.59	0.00	11.03	2.05	3.39	0.00	0.00	4.10	2.77	3.91	0.00	5.95	11.97	0.00	3.43
TC 50	320.00 >	1000.00 >	897.00 3200.00	1000.00 >	18.50	320.00 > 585.00	320.00 > 692.00	320.00 > 608.00	210.00 252.00	320.00 > 835.00	320.00	5.77	320.00 > 767.00 320.00	234.00	320.00 710.00 320.00	320.00 > 213.00
10 50	320.00 > 0.00 > 1	631.00 > 1	204.00	0.00 > 1	9.00	94.30 >	203.00 > 0.00	173.00 >	51.30	115.00 > 203.00	0.00 > 255.00 > 1	3.10	296.00 × 129.00 0.00 ×	19.50	320.00	93.30 >
A1 25	0.00	2.91	3.9	24.88	2.39	7.04	2.30	0.58	33.13	2.35 5.35	3.10	2.52	1.87 9.06 1.76	6.09	3.31	8.13
55 21	320.00 >	1000.0 > 2630.0	527.00 1630.0	1000.0 > 728.00	11.70	320.00 > 378.00	320.00 > 371.00	68.30	155.00	100.00	253.00	3.48	210.00 544.00 320.00 >	76.60	283.00 502.00 320.00	320.00 > 150.00
10 25	% 0.00	344.00 >	132.00	40.20 >	00.00	45.40 >	139.00 > 305.00	118.00	9.00	42.50	81.60	1.38	112.00 60.00 182.00 ×	12.60	100.00 152.00 0.00 ×	39.40 >
oiff.	0.289	0.792	0.823	0.902	0.576	0.630	0.714	0.636	0.669	0.339	0.654	0.716	0.646	0.658	0.733	0.718
14	₹	VBR X54	VCO XS7	VC1	VC3	XX	X S	XXXXXX	VX	UNB VX2	XXX XXX	XH9	XXK YRB	XXX	XXM	XK2
Test	03/15/90 UNG 05/03/90 VAN	04/18/90 VBR 06/26/90 X54	04/18/90	04/18/90	04/18/90	06/05/90 07/12/90	06/05/90	06/05/90	03/08/90 UFO 05/03/90 VX1	03/15/90	06/05/90	06/12/90 07/12/90	06/12/90 08/02/90 09/11/90	06/12/90	06/15/90 WSY 08/02/90 XXH 09/11/90 YRE	07/17/90
Ship- ment	33	33	33	33	33	29	19	79	33	33	19	19	222	22	222	69
AVS S No. m	6583	6584 6584	1099	6603	6607 6607	6714 6714	6722	6730	6745	8748	6753	6792	6828 6828 6828	6837	6861 6861 6861	6873

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ig.	Assay Type	ĔĔ	EE	FF	ĔĔ	EE	ĦĦ	FEE	E E	FFF	E E	I I	I I	FF	ĒĒ	H H	THE	
	Ĭ	14.78	12.37	2.39	36.84	32.49	36.17	11.78 12.85 0.00	0.00	8.51 19.33 4.99	16.74 0.58	15.06	7.06	23.20	19.93	3.89	1.21	
	ïs	1.44 > 0.00 >	2.02 > 0.00 >	1.07 >	21.45 > 0.00	5.28 >	11.80 > 0.26	3.35	0.00	0.00 5.37 1.17	1.86 >	1.95 > 2.46	1.71 >	4.77 > 3.53 >	2.22 > 0.40	0.00	0.00 >	
	A1 95	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	3.17	0.00	0.00	0.0	
	7C 95	320.00 935.00	96.60	320.00	320.00	320.00	320.00 923.00	303.00 305.00 291.00	309.00	309.00 309.00 305.00	320.00	320.00	320.00	320.00	320.00 958.00	320.00	320.00	
	36 31	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	316.00 >	0.00	0.00	0.00	
	AI 50	1.44	2.72	1.07	21.45	5.28	19.44	0.00	0.00	0.00	0.00	1.95	1.71 2.39	5.27	2.22	0.00	0.00	
	TC 50	320.00 >	66.00	320.00 > 806.00	320.00 > 817.00	320.00 > 585.00	320.00 > 269.00	151.00 167.00 82.30	210.00	210.00 210.00 171.00	320.00 >	320.00 >	320.00 > 732.00	320.00 > 927.00	320.00 > 479.00	320.00 550.00	320.00	
	10 50	222.00 > 0.00	24.30	299.00 > 0.00	14.90 >	60.60 > 75.20	16.50 > 320.00	0.00 27.70 0.00	0.00	0.00 28.90 82.70	172.00 >	164.00 >	187.00 > 306.00	60.70 >	144.00 > 179.00	308.00	0.00 > 892.00 >	
	AI 25	3.87	70.0	1.87	90.99	69.60	194.29	5.24 6.14 0.00	0.00	2.90 12.62 2.12	5.31	3.87	2.34	6.57	8.27	5.28	1.79	
	TC 25	320.00 > 202.00	30.40	320.00 > 502.00	320.00 > 490.00	320.00 > 378.00	194.00 > 83.00	80.60 92.70 45.30	155.00	155.00 155.00 97.00	320.00 >	320.00 >	320.00 > 454.00	290.00	320.00 > 71.40	253.00 276.00	320.00 > 563.00	
	10 25	82.70 > 0.00	12.10	171.00 × 881.00	4.84 > 706.00	6.45 > 43.50	1.00	15.40 15.10 0.00	0.00	53.40 12.30 45.70	60.30 >	70.60 > 127.00	137.00 >	44.10	38.70 > 60.90	47.90	179.00 > 505.00	
	oiff.	0.766	0.718	0.717	0.760	0.760	0.742 < 1.028	0.782 0.607 1.011	0.837	0.837 0.607 0.958	1.078	0.858	0.810	0.842	0.725	0.821	0.750	
	*	XNG	XNH	XR3	XRO	XRD	XU3	XXX XXX YT8	X1N YT8	XXX YTS	x10	X10 YU1	X YOU	X1T	XU6	XU7	XUA YN2	
	Test Date	07/19/90 XNG 09/05/90 YKK	07/19/90	07/24/90	07/24/90	07/24/90	07/26/90	06/21/90 08/02/90 09/13/90	06/21/90 09/13/90	06/21/90 08/02/90 09/13/90	06/21/90 09/13/90	06/21/90 09/13/90	06/21/90 09/13/90	06/21/90 09/13/90	07/26/90	07/26/90	07/26/90 XUA 09/06/90 YN2	
	ment	69	69	69	69	69	69	222	2 2	222	28	2 2	2 2	22	69	69	69	
	NO.	6903	7069	6923	2769	6943	945	0769 0769 0769	9769	7769 7769 7769	8769	6269	6983	8869	7003	7007	7011	

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Assay Type	FF	ĒĒ	ĒĒ	ĒĒ	ĒĒ	ĒĒ	ĒĒ	ĒĒ	ĔĒ	ĒĒ	ĒĒ				
¥	0.00	3.14	4.33	24.46	13.42	14.69	8.47	13.81	4.58	0.70 6.56	7.94	2.21 1.04 14.18	2.40 11.05 17.70	4.33	5.25 12.05 14.09
15	0.00	3.00	1.05 > 2.82 >	2.41 >	2.79	2.18 > 0.95 >	2.48	2.15	1.53	0.00	1.20 >	0.00 \$ 2.46 \$	0.00	0.00	0.00 > 0.00 > 2.27 >
A1 95	9.0	3.22	0.00 \	1.09 v	0.00	1.07 > 0.00	0.00	0.0	0.00	0.00	0.00	0.00	0000	988	888
7C 95	320.00	308.00	320.00	320.00 > 967.00	320.00 921.00	320.00 >	320.00 949.00	309.00 821.00	307.00	305.00	320.00 1000.00	309.00	100.00 959.00 320.00	100.00 923.00 959.00	320.00 955.00 965.00
10 95	00.00	97.30	0.00 >	293.00 >	90.0	300.00 >	0.00	0.0	0.00	0.00	0.00	0000	0.00	.000 000 000	00.00
A1 50	0.00	0.00	1.05	2.41	3.95	2.18	4.27	2.94	2.27	0.00	1.20	0.00	2.67	0.00	0.00 0.00 3.13
10 50	320.00	168.00	320.00 >	320.00 >	241.00 289.00	320.00 >	305.00	208.00	194.00	169.00 165.00	320.00 >	210.00 181.00 210.00	100.00 590.00 320.00 >	100.00 296.00 495.00	320.00 553.00 646.00
10 50	0.00 \$	0.00	306.00 >	133.00 > 93.40	61.00	147.00 > 938.00 >	71.40	0.00	85.50	0.00	266.00 > 601.00 >	0.00	0.00 221.00 72.90 4	0.00 \$ 246.00	0.00 0.00 206.00
AI 25	1.01	1.89	3.20	15.18	67.4	3.34	3.70	3.19	2.49	1.27	3.84	0.00	3.38	0.00	1.42 3.29 3.93
TC 25	77.30	84.30 156.00	320.00 >	320.00 >	171.00	320.00 > 887.00	177.00 267.00	152.00 154.00	130.00	91.50 90.60	320.00 >	155.00 111.00 155.00	386.00 258.00	100.00 198.00 232.00	320.00 > 329.00 469.00
10 25	0.00	36.80	100.00 >	21.10 >	38.00	95.70 > 548.00	47.80	00.0	52.30	72.20	83.30 >	0.00	0.00 > 114.00 47.10	0.00	225.00 × 100.00 119.00
Diff.	0.690	0.678	0.609	0.501	0.553	0.520	0.520	0.503	0.676	0.676	0.686	0.573	0.502 0.942 0.908	0.655 0.987 1.089	0.655 0.987 0.783
= *	X X	XUG	XX3	XX7	XX8	XXB YR4	XXB	XXC	XXH YR7	XXH YR7	XXI	YDN 272 20F	750 272 295	YFL 280 29.3	7FL 280 20K
Test	07/26/90 XUD 09/06/90 YN4	07/26/90	07/31/90 09/06/90	07/31/90 09/06/90	07/31/90 09/11/90	07/31/90 09/11/90	07/31/90 09/11/90	07/31/90 09/11/90	08/02/90	08/02/90 09/11/90	08/02/90	08/29/90 10/02/90 10/25/90	08/29/90 10/02/90 10/25/90	08/30/90 10/02/90 10/25/90	08/30/90 YFL 10/02/90 280 10/25/90 29K
Ship- ment	99	99	99	99	69	69	69	% %	69	69	66	222	222	222	222
AVS S	7017	7022	7023	7032 7032	7033	7039	7040	7042	7047	7048	7049	7055 7055 7055	7057 7057 7057	707 1707 1707	2072 2707 2707

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Acces	Type	FFF	EEE	EEE	FEE		EEE	EEE	FFF			E E	EE	FE	
	Ĭ	3.70	2.34 12.67 29.69	6.82 23.51 42.40	6.23 0.77 21.13	1.88 2.17 17.33	2.91 10.96 33.38	7.04 31.69 36.77	6.81 7.45 13.64	10.83	28.98 1.10 25.87	2.04	3.35	22.45 11.58	
	ï	0.00	0.00	0.92 > 1.47 > 12.88 >	0.00 > 3.34 >	3.16 ×	0.00	0.00 > 6.86 > 10.67 >	0.00	9.53	0.00 78.8	4.20	0.00	3.93 >	
	A1 95	0.00	3.68	0.00	0.00	96.6	0.00	0.00	0.00	0.00	88 88	0.0	0.00	0.00	
	TC 95	320.00 1000.00 1000.00	300.00 305.00 315.00	100.00 97.10 95.80	93.40 262.00 100.00	78.80 100.00 93.00	100.00	320.00 1000.00 1000.00 >	320.00 800.00 924.00		00.0001	320.00	307.00	1000.00	
	10 %	0.00	0.00 0.00 85.70	0.00 0.00 9.31	0.00	0.00	0.00	0.00 > 0.00 > 1030.0 >	0.00	90.0	88 88	0.00	0.00	0.00	
	AI 50	0.00	0.00 2.26 8.73	2.98 2.66 24.01	0.00	0.00	0.00 26.09	0.00 6.86 10.67	0.00	0.00	0.00	5.95	0.00	0.00	
	TC 50	320.00 507.00 660.00	124.00 171.00 208.00	27.00 53.40 47.90	25.80 57.00 64.60	16.40 28.20 28.30	29.90 51.40 94.30	320.00 1000.00 1000.00	233.00 233.00 269.00	1000.00	1000.00 1000.00 985.00	320.00	186.00	1000.00 >	
	10 50	0.00 0.00 320.00	0.00 75.80 23.80	9.07 20.10 2.00	0.00	0.00 6.00 8.00	0.00	0.00 > 146.00 > 93.80 >	0.00 63.80	0.00 \$1.90	60.60 v	0.00 >	320.00	218.00 \$	
	AI 25	0.00	1.00	2.07 19.61 30.04	0.00	1.01	1.68 4.49 13.69	15.77	2.97	7.92	20.22	0.00	1.34	32.22 5.95	
	10 25	253.00 402.00	52.10 97.40 148.00	8.30 29.50 25.70	16.60 7.11 41.30	7.66 18.70 19.20	7.09 22.50 24.10	320.00	150.00 166.00 171.00	781.00	220.00 606.00 616.00	320.00	119.00	854.00 851.00	
	10 25	0.00 0.00	52.10 35.10 13.50	4.02 1.50 0.86	0.00 17.90 5.56	7.59	4.21 5.01 1.76	211.00 > 63.40 > 24.80 >	\$0.50 88.10 41.50	113.00	28. 28. 28. 28. 29. 28.	0.00	88.50 152.00	26.50	
	Diff.	0.700 1.051 0.853	0.751 0.976 0.853	0.751	0.811 0.975 0.796	0.811 0.975 0.808	0.640 0.976 0.808	0.966	0.716 0.966 0.999	0.651	1.027	0.592	0.421	0.935	
•	*	281 281 20 29L	00 YFR 00 282 00 20L	7 YFR 20 282 20 294	70 YFS 70 Z83 70 Z9M	7 YFS 70 283 70 29N	20 YFT	7 7KA 20 285 20 290	00 YKB 00 285 00 290	00 Y28	0 Y1C 0 Y1C 0 Y1D 0 Y1D	0 Y2H 0 Y1D	00 Y4K 00 YTG	00 Y6M	
1001	Date	08/30/90 10/02/90 10/25/90	08/30/90 10/02/90 10/25/90	08/30/90 10/02/90 10/25/90	08/30/90 10/02/90 10/25/90	08/30/90 10/02/90 10/25/90	08/30/90 10/02/90 10/25/90	09/05/90 10/02/90 10/25/90	09/05/90 10/02/90 10/25/90	08/10/90	09/13/90 09/13/90 08/10/90	08/10/90 09/13/90	08/14/90	08/16/90	
Shin	ment	222	222	222	222	222	222	222	222	22 8	28 88	22	22	22	
		7074	7083 7083	7084 7084 7084	7085 7085 7085	7086 7086 7086	7087 7087	7092 7092 7092	7094 7094 7094	7116	2002	7308	7320	376	

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Assay Type	T T T	ĔĒ	EE	FF	ĒĒ	T I	ĔĔ	T T	T I	ĒĒ	ĒĒ	EE	FFFF	FE	ËË	E E	
141	2.28	1.23	4.26	5.19	1.48	2.68	5.16 0.18	1.20	2.54	2.86	0.85	5.36	3.20	2.16	14.40	10.97	
3	0.00 v	0.74 >	1.43 > 2.09 >	0.14 > 0.00 >	0.47 >	1.07 > 0.00	1.42	0.00	0.00	0.25	0.69 \$	1.29 \$ 0.00	0.88 0.75 0.00 0.00	1.00.0	3.16	1.88 v	
A1 95	0.0	0.0	9.00	90.0	88	90.0	99.	88	9.0	0.00	99.	0.00	0.000	9.00	9.0	0.00	
7c %	1000.00	1000.00 3200.00	320.00 961.00	320.00 917.00	320.00 1000.00	320.00 972.00	306.00	1000.00 3200.00	320.00 320.00	1000.00 2870.00	1000.00 3110.00	320.00 1000.00	320.00 944.00 320.00 1000.00	320.00 1000.00	1000.00	320.00	
10 95	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.0000	0.00	0.00	0.00	
A1 50	9.0	0.0	3.05	3.52	1.0	0.00	2.24	1.07	2.31	2.71	0.00	1.29	1.38	0.0	0.00	0.00	
10 50	1000.00 > 251.00	1000.00 >	320.00 > 614.00	320.00 >	320.00 >	320.00 >	184.00 145.00	1000.00 >	185.00 320.00	703.00	1000.00 >	320.00 > 735.00	320.00 × 443.00 320.00 × 639.00	320.00 >	986.00 974.00	320.00 >	
10 50	930.00 >	919.00 \$	223.00 > 201.00	%.% 0.00	290.00 > 919.00 >	245.00 >	82.20	938.00 >	0.00	259.00 188.00	978.00 >	247.00 > 0.00	248.00 × 320.00 × 305.00 × 0.00	320.00 >	202.00	170.00 >	
AI 25	0.00	1.26	3.17	1.12	0.88	1.72	2.31	1.33	1.00	3.93	1.21	2.05	1.62	1.83	4.76	2.72	
TC 25 A	876.00 106.00	683.00	. 320.00 > 420.00	12.90	137.00	261.00 329.00	116.00	728.00	32.00	66.00 516.00	676.00 1060.0	> 320.00 > 527.00	217.00 241.00 320.00 >	. 320.00 > 563.00	639.00	> 320.00 > 505.00	
10 25	507.00	542.00	148.00 >	3.20	156.00 542.00	152.00	50.30	548.00 935.00	45.80	161.00	559.00 859.00	0.00	151.00 149.00 171.00 v	175.00 \$	134.00	118.00 0	
biff.	0.935	0.949	0.950	0.872	0.953	0.873	0.873	0.853	0.994	0.934	0.832	0.873	0.964 1.038 0.882 1.129	0.899	0.800	0.800	
* 5	00 Y W	79X 00	00 Y80 YVX 00	00 Y8P	O YBR	20 Y8T	20 Y8T	20 Y8U 20 236	20 Y8W	20 YAY 20 238	20 YB0	00 YB1	782 782 70 783 70 783 70 783 70 783	20 YB4	20 YB6 23 Z3C	00 YB6	
Test Date	08/16/90 Y6M 09/18/90 YVR	08/16/90 Y6V 09/18/90 YV4	08/21/90 09/18/90	08/21/90 09/18/90	08/21/90 09/18/90	08/21/90 09/27/90	08/21/90	08/21/90 09/27/90	08/21/90 09/27/90	08/23/90	08/23/90 09/27/90	08/23/90 09/27/90	08/23/90 09/27/90 08/23/90 09/27/90	08/23/90 09/27/90	08/23/90 09/27/90	08/23/90 09/27/90	
Ship- ment	22	22	22	22	22	22	22	22	22	22	22	22	2222	22	22	22	
AVS S	7350	7368	23.25 27.27	73.75 73.75	7379 7379	7383	7385	7386	7391	73%	7403	7405	2406 7408 7408 7408	7410	7414	7415	
~ ~	1010	1-1-	1-1-	1-1-	1-10	1-1-	1-1-	1-1-		1-1-	1-1-	1-1-	1-1-1-1-	1-1-	1-1-	1-1-	

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Assay		ĒĒ	ĒĒ	ĒĒ	ĒĒ	ĒĒ	H H	ĔĔ	ĒĒ	ĒĒ	EE	FE	FF	ĒĒ	EEEE	ĒĒ	
TAI	3.20	23.88	5.85	0.00	7.17	0.00	0.00	18.38 5.52	7.86	9.69	3.98	7.34	9.51	0.25 32.26	2.83 0.00 9.51	0.00	
20	1.26 > 0.00	4.02 >	1.73 >	0.00	2.34 >	2.96	1.82 ×	1.63 >	1.52 > 2.06	0.9%	0.00 >	1.08	2.09 >	9.05	0.00 0.33 2.24 0.25	0.00	
8 8	0.00	1.16 > 0.00	 	80.0	88.	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	3.59	8888	88.	
70 % 21	320.00 962.00	320.00 > 973.00	320.00 958.00	320.00	320.00 964.00	979.00	1000.00	306.00 320.00	320.00	320.00	320.00	320.00	320.00	3200.00	320.00 3200.00 320.00 3200.00	320.00	
8	00.00	277.00 > 0.00	0.00	0.00	0.00	0.0	3010.0 > 3	0.00	0.00	0.00	0.00	0.00	0.00	0.00 > 3	0.0000	0.00	
AI 50	1.26	6.02	1.73	0.00	3.24	0.00	0.00	2.58	1.52 3.25	1.37	3.08	1.02	0.00	0.00	0.00 1.32 0.00 3.49	3.88	
10 50	320.00 >	320.00 >	320.00 × 575.00	320.00	320.00	516.00 638.00	683.00	184.00	320.00 > 603.00	263.00	320.00	320.00 > 622.00	320.00	320.00	320.00 3200.00 > 320.00 2410.00	320.00	
10 50	254.00 > 0.00	73.60 0.00 v	185.00 > 0.00	0.00 >	0.00 >	0.00	0.00	71.30	211.00 > 185.00	192.00	0.00 > 235.00	312.00 > 212.00	370.00 > 1	320.00 > 3	2430.00 > 3 0.00 > 3 691.00 2	0.00 > 157.00	
AI 25	2.41	7.10	3.12 2.13	3.19	2.17	9.00	0.00	44.62	2.20	1.71	2.15	1.56	2.03	18.09	0.00 1.70 3.30	0.00	
10 25	320.00 >	320.00 >	320.00 >	320.00	320.00 >	251.00	390.00	116.00	320.00 >	180.00	215.00	267.00	320.00 >	320.00	155.00 1920.0 160.00 1550.0	320.00	
10 25	133.00 >	45.00 >	102.00 >	0.00 >	147.00 >	0.00	1300.0	2.61	145.00 >	105.00	100.00	170.00	158.00 > 39.50	160.00	0.00 1130.0 0.00 470.00	0.00 >	
Diff.	0.934	0.828	0.888	0.513	0.595	1.235	1.152	0.557	0.893	1.089	1.192	1.095	1.0%	1.085	1.123 0.943 0.950 0.950	0.849	
± *	08/23/90 YB7 09/27/90 23E	08/23/90 YBB 09/27/90 23E	/90 YBB /90 23F	/90 YDH	/90 YDJ	/90 YXS	09/20/90 YXV 10/25/90 200	/90 XX2 /90 YRA	10/30/90 YZY 10/30/90 ZVM	/90 205	902 06/ 902 06/	09/25/90 207 10/30/90 2VP	09/25/90 207 10/30/90 2vg	10/11/90 2FV 11/06/90 110	/90 2JL /90 13M /90 130	/90 2JQ /90 13R	
. Test Date	08/23	08/23	08/23/90	08/29/90	08/29/90	09/20/90	09/20/90 10/25/90	07/31/90	09/25/90	09/25/90 10/30/90	09/25/90	69/25 10/30	09/25 10/30	10/11/90	10/16/90 11/08/90 10/16/90 11/08/90	10/16/90	
Ship-	22	22	22	22	22	RR	RR	22	RR	RR	22	RR	RR	% K	* 5 7 7 7	22	
AVS.	7417	7418 7418	7424	7430	77.77	7439	7445	6772	6972	7872	7485	7487	7488	3010	255 255 255 255 255 255 255 255 255 255	7935 7935	

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Assay Type	E E	ĒĒ	ĒĒ	ĒĒ	ĔĒ	EE	
¥	0.00	1.10	9.40	0.52	0.0	8.59	
15	5.50 ×	0.00 >	0.00 \$	0.00	0.18	1.21 > 0.00 >	
A1 95	3.03	88.	88	88	88	88	
25 21	320.00	320.00	320.00	320.00	940.00	3020.00	
20 21	320.00	0.00	0.00	9.00	88	8.8	
AI 50	0.00	0.00	3.73	1.84	77.0	2.52	
TC 50	320.00	218.00	320.00	320.00 1570.00	29.70	958.00	
10 50	7.8	0.00	0.00 >	0.00 \$	919.00	380.00	
AI 25	8.3	3.64	0.00	0.00	0.30	3.04	
10 25	251.00	159.00	91.50	\$20.00	162.00	459.00	
52 21	0.00	9.02	0.00 \$	387.00	542.00	151.00	
Diff.	1.042	1.001	0.920	0.920	0.908	0.910	
=	727	13U	23V 13V	14	19H	272	
Test	16/90/11 11/06/90/11	10/16/90 ZJU 11/08/90 13U	10/16/90 2JV 11/08/90 13V	10/16/90 2JV 12/04/90 1M4	11/15/90 19H 12/04/90 1MH	11/15/90 190 12/21/90 277	
Ship-	22	22	22	22	ĸĸ	ĸĸ	
AVS .	0362	7942	776	7945	78.2	8212 8212	

This value is a virus rating (VR) rather than a TAI. The VR is a measurement of selective antiviral activity that takes into account the degree of inhibition of virus-induced CPE and the degree of cytotoxicity produced by the test compound similar to IAI. IAI is more accurate with MII measurements. The differential is the difference in the cell control and the virus control optical densities. DIFRNTL =

(Viral) inhibitory concentration 25%, 50% and 95% = The drug concentration (µg/ml) that inhibited viral CPE by 25%, 50% or 95% calculated by using a regression analysis for semilog curve fitting. IC25.50.95

Antiviral Index = A single point ratio of the antiviral and anticellular effect of the compound, calculated with 25%, 50% or 95% reduction values (Cell) toxicity concentration 25%. 50% and 95% a The drug concentration (µg/ml) that reduced cell viability by 25%, 50% or 95%. (calculated by dividing the TC_{25,50,95} by the IC_{25,50,95}). TC25,50.95 AI 25,50,95

Selectivity Index = A ratio calculated by dividing the IC₂₅ by the IC₅₀ (based upon 6 one-half-log₁₀ dilutions, µg/ml, the maximum scale is 0-320).

lotal Antiviral Index st The area between the cytotoxicity and the antiviral curves (based upon a scale of 0-100%).

TAI ACT

SI

Activity = A *** denotes a test that produced >25% reduction in CPE. A *** denotes an inactive test (i.e. <25% reduction in CPE).

7/12/91 JJK/ADB/ap

4.1.7 Sandfly Fever Virus (SF):

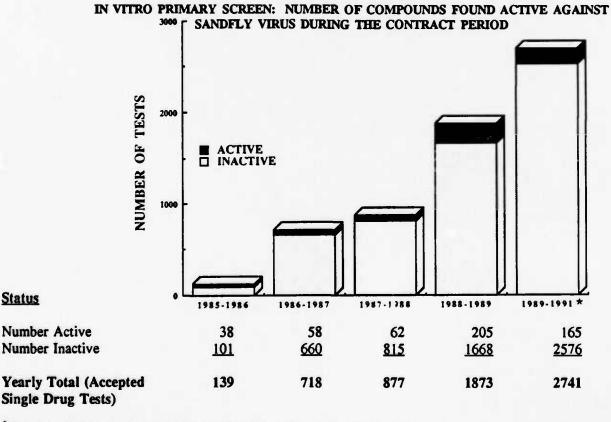
The number of single drug tests carried out against SF during this contract period is summarized in yearly increments in Figure 36. During this five-year period two main in vitro antiviral assay protocols were implemented:

- 1. A standard CPE inhibition assay by virus rating (VR) (Annual Report, December 15, 1988, Section 3.2.4).
- 2. Since November, 1988, MTT based-antiviral assay format.

A total of 7833 tests were performed during this contract period using both assay types. Routine testing was changed to the MTT-assay format to improve the efficiency and quality of the primary screening program in addition to being more cost-effective. Ribavirin (AVS-0001) was tested in each standard virus rating (VR) CPE-inhibition assay as a positive control compound. Results of these positive controls (VR tests) were used as a guideline to assess the quality of each assay.

After the testing was converted to the MTT-assay format, we performed a total of 414 control compound assays with Ribavirin during the last 26 months of the contract period. During this time 610 tests were internal (+++) virus load, cell load, and other quality control tests. Three hundred twenty-two (322) tests were considered unsatisfactory based on the criteria of the quality controls set during this reporting period. The rest, totaling 4614 were actual single drug MTT-assays. The total number of MTT-assays (5960) tested during the last two years represents a 218% increase (improvement) in the total testing output as compared to the total of 1873 tests performed during the first 3 years of this contract.

Out of the 6487 accepted single drug tests, 566 compounds demonstrated antiviral activity at greater than 50% reduction levels. This represents around 9% of the tested compounds having *in vitro* antiviral activity against SF-virus. The remainder, 5921 compounds (91%), were considered inactive with both assay protocols (Figure 36).



Represents 14-month period (November 15, 1989 - January 31, 1991)
Figure 36

Five-Year

528

5820

6348

Totals .

- 4.1.7.1 <u>SF-Quality Controls:</u> Two positive control compounds (Ribavirin and 2-Thio-6 Azauridine) were used in the daily assay sets as antiviral activity quality controls. The antiviral performance of the unknown compounds is compared to that of the positive control compounds. Compounds with equal to of better antiviral potency are considered active and are worthy of further *in vitro* profile studies and *in vivo* testing.
- 4.1.7.1.1 Antiviral Activity of Ribavirin vs SF Virus: A summary of the antiviral and cytotoxicity performance of the primary control compound, AVS-0001 (Ribavirin) is presented in Figure 37-A for 226 tests performed during November, 1989 through January, 1991.

Control Compound-Antiviral Performance: Ribavirin (AVS-0001) has been the sole control compound against SF in these MTT-assay screens. The mean and median antiviral inhibition and cytotoxicity patterns of the positive control drug (Ribavirin) are illustrated in Figure 37-A.

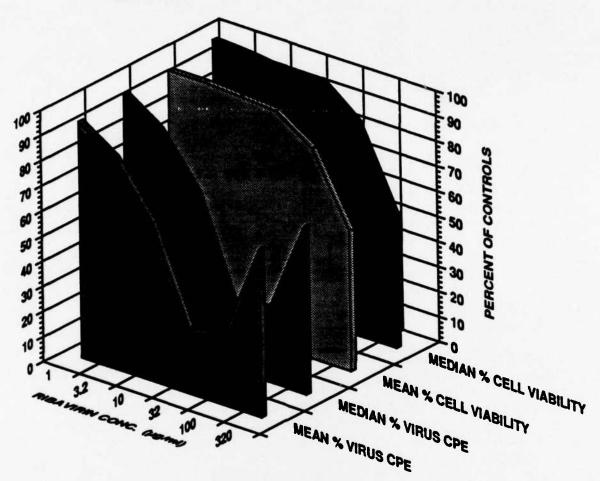
The 226 control tests performed with Ribavirin gave a mean Total Antiviral Index (TAI) of 33.5% (SD \pm 13.93) and the median value was 31.28%. The TAI measures the overall antiviral effectiveness of the compound and it ranged from \sim 0.80-73% during this period. The mean Selectivity Index (SI) was 15.33 (SD \pm 15.89) and the median SI value was 10.79, indicating moderate antiviral selectivity for Ribavirin against SF virus. SI ranged from \sim 0-109 during this period. However, the closeness of the mean and median values indicate that the present execution of the SOP is consistent and repeatable.

The mean Antiviral Index 25% (AI₂₅) value was 40.9 (SD \pm 47.5). The median AI₂₅ value was 25.5 (range 0 - 320). The mean Antiviral Index 50% (AI₅₀) was 24.38 (SD \pm 17.9) with a median of 19.21 (range 0 - >109). The mean Antiviral Index 95% (AI₉₅) was 2.2 (SD \pm 5.0), with a median of 0 (range 0 - 36.22). This indicates that the control compound, Ribavirin, does not consistently reach 95% antiviral reduction levels.

The mean Antiviral Inhibitory Concentration 25% (IC₂₅) was 7.37 μ g/ml (SD \pm 4.95). The median IC₂₅ value was 5.72 μ g/ml (range = 0 - 30.0 μ g/ml). The mean Antiviral Inhibitory Concentration 50% (IC₅₀) was 17.76 μ g/ml (SD \pm 11.81). The median IC₅₀ value was 15.65 μ g/ml (range = 0 - 79.6 μ g/ml). The mean Antiviral Inhibitory Concentration 95% (IC₉₅) was 9.81 μ g/ml (SD \pm 21.66). The median IC₉₅ value was 0 μ g/ml (range from 0 - 94.5 μ g/ml). This discrepancy indicates that the control compound Ribavirin does not consistently reach 95% reduction levels. During this reporting period, the highest starting concentration of Ribavirin (100 μ g/ml) was increased from previous high dose of 100 to 320 μ g/ml to properly evaluate the maximum antiviral effect of Ribavirin.

The average maximum antiviral inhibitory level of 226 Ribavirin tests (Figure 37-A) was reached at 32 μ g/ml of the compound with 75% antiviral effect. Maximum antiviral effect (~80%) was found with a simultaneous ~5% cytotoxic suppression. Above (100 μ g/ml) concentration Ribavirin starts to lose its antiviral potency (~30%) at 320 μ g/ml while simultaneously the Ribavirin becomes maximally toxic (~45%).

RIBAVIRIN - VS - SF VIRUS



CONCENTRATION (µg/ml)

%	Viral	CPE	% Cell Viability

Conc.(rg/ml)	1	3.2	10	32	100	320	1	3.2	10	32	100	320
Mean	94	84	63	26	29	67	96	96	95	94	84	56
Median	96	86	66	26	30	67	99	98	96	97	85	53
Std. Dev.	0.07	.11	0.20	0.18	0.19	0.18	0.06	0.05	0.05	0.07	0.14	0.16

Figure 37-A

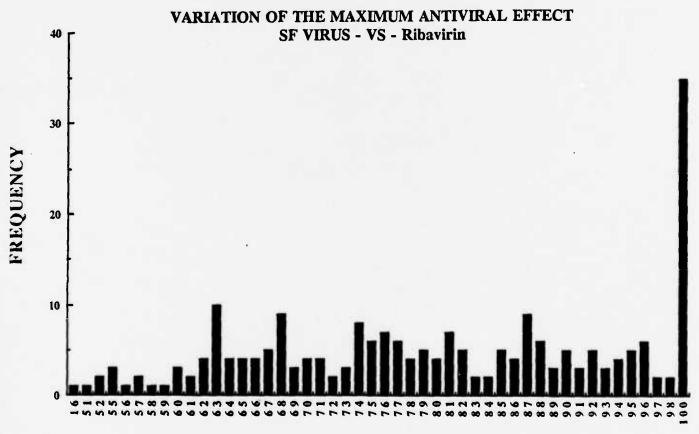
Average Antiviral and Cytotoxicity Values for 226 Positive Control Compound Tests

4.1.7.1.2 <u>Maximum Antiviral Effect of Ribavirin vs SF Virus:</u> Since the metabolic activity of the cells was an unknown function during the testing period, it was monitored indirectly by measuring the maximum antiviral effect of the control compound Ribavirin. This demonstrated the amount of infectious virus that was produced by the cells (Maximum Percent CPE).

A bar graph scatter plot (Figure 38-A) depicts the distribution of the maximum antiviral reduction values of all 226 control compound assays for Ribavirin. The results indicate that the average maximum antiviral reduction obtained with the present SOP is around 80% (SD \pm 14.30) reduction levels. The maximum reduction levels vary from 16 - 100% but remain quite consistently around the median of 81%. The assay control values give a shifted half-bell-shaped distribution curve toward the maximum 100% reduction level. This indicates quite a consistent day-to-day performance of the control compound in the SF-MTT assay.

During this period the positive control compound performance criteria for Ribavirin versus the SF virus was set at 50% reduction level. All assays in which Ribavirin did not meet this accepted quality control level ($\geq 50\%$) were rejected (i.e., 322 unsatisfactory tests).

Ribavirin is active in vitro against SF virus and functions as a reasonable quality control compound. On the other hand, regardless of the performance of the SF-quality control drug Ribavirin, around 165 other compounds have equal or better antiviral activity against SF virus than AVS-0001. (See 95% and 50% reduction summaries).



PERCENT CPE REDUCTION

Figure 38-A

Maximum Antiviral CPE Reduction (%).

Summary of 226 Control Tests.

4.1.7.1.3 Cellular Cytotoxicity of Ribavirin vs SF Virus:

SF-Control Compound-Cytotoxicity Performance: The 226 cytotoxicity values of the positive control compound Ribavirin are also very consistent. The mean cell Toxic Concentration 25% (TC₂₅) was 176.52 μ g/ml (SD \pm 94.76) and the median was 165 μ g/ml (range of 0.78 - 320 μ g/ml). The mean cell Toxic Concentration 50% (TC₅₀) value was 290.7 μ g/ml (SD \pm 48.00) and the median was 320 μ g/ml (range of 96-320 μ g/ml). The mean cell Toxic Concentration 95% (TC₉₅) value was 320 (SD \pm 0) and the median was 320 (range of 320-320 μ g/ml). This discrepancy indicates that the control compound Ribavirin does not consistently reach 95% cytotoxicity levels.

As can be seen from Figure 37-A, the toxicity starts to become measurable above the concentration of 100 μ g/ml and the maximum toxicity has not been reached at 320 μ g/ml. Further increase of the concentration of Ribavirin would be needed to properly evaluate the maximum cytotoxicity of Ribavirin.

Also, Figure 37-A indicates that when the cytotoxicity reaches $\sim 10\%$ at $100~\mu g/ml$, the control compound (Ribavirin) has reached simultaneously its maximum antiviral effect (80%). The cytotoxic effect of Ribavirin is insignificant between 1 and 100 $\mu g/ml$. The average cytotoxicity reached $\sim 45\%$ at 320 $\mu g/ml$, which is the highest Ribavirin concentration tested.

Ribavirin has a definite cytotoxic suppression on cellular metabolism and growth. However, the TC_{25} and TC_{50} toxicity could not be consistently achieved with the 100 μ g/ml concentration of Ribavirin. Therefore, a readjustment to 320 μ g/ml (as being the highest Ribavirin concentration tested) was done during this reporting period. However, at this concentration (320 μ g/ml) the TC_{50} and TC_{95} cannot yet be measured consistently.

4.1.7.1.4 SF-Assay Plate Quality Controls: Cell Load and Virus Load Parameters (Ribavirin): The MTT assay is fundamentally dependent upon the quality of the assay plates. Our large-scale antiviral testing is dependent upon the uniformity of the test plates produced for the daily assays. Equal numbers of cell load and virus load as well as the consistent performance of the reagents used daily was monitored. A sample of the plate variation controls for the period of November, 1989 through January, 1991, is presented in Figures 39-A, 40-A and 41-A.

SF-Control Compound-Cell Load Performance: A bar graph scatter plot of the mean cell control (O.D. reading) of 226 control assays is plotted in Figure 39-A. The results indicate that the cell O.D. readings reached a mean of 1.030 (SD \pm 0.160) with a median of 1.010 (range of 0.681 - 1.566). This indicates that a uniform and equal number (18,000 cells/well) of cells are being loaded into every well in the 96-well plate during the day-to-day operation. The cells reduced MTT to formazan giving maximum blue color uniformly and consistently.

SF-Control Compound-Virus Load Performance: A bar graph scatter plot of the mean virus load O.D. readings of the 226 control assays is presented in Figure 40-A. The results indicate that the average virus load O.D. reading reached a mean of 0.250 (SD \pm 0.110 with a median of 0.240 (range of 0.018 -0.646). This demonstrates that a good cell destruction is taking place and a uniform load of virus (32 TCID₅₀) is administered on the cell monolayer with very consistent viral CPE results.

SF-Control Compound-Assay Differential Performance: A bar graph scatter plot of the mean O.D. differential values of the 226 control assays is provided in Figure 41-A. The results indicate that the average differential O.D. reading is 0.780 (SD \pm 0.162) with a median of 0.763 (range 0.480 - 1.456). The single bell-shaped curve is reasonably sharp and uniform. This reflects that the assays are executed consistently and are repeatable during day-to-day operation with close to 78% measurement accuracy.

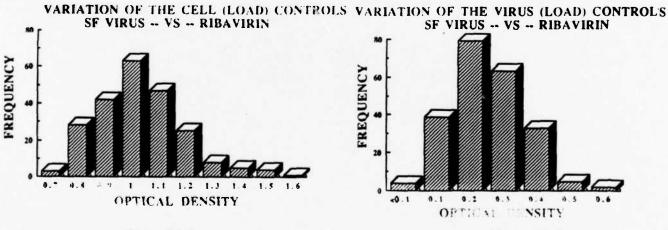
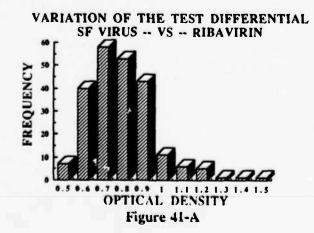


Figure 39-A

Figure 40-A



4.1.7.1.1 Antiviral Activity of AVS-6724 (2-Thio-6-Azauridine) vs SF Virus: A summary of the antiviral and cytotoxicity performance of the second control compound, AVS-6724 (2-Thio-6-Azauridine) is presented in Figure 37-B for 42 tests performed during November, 1989 through January, 1991.

Control Compound-Antiviral Performance: 2-Thio-6-Azauridine (AVS-6724) has been tested as a possible control compound against SF in these MTT-assay screens. The mean and median antiviral inhibition and cytotoxicity patterns of this second positive control drug are illustrated in Figure 37-B.

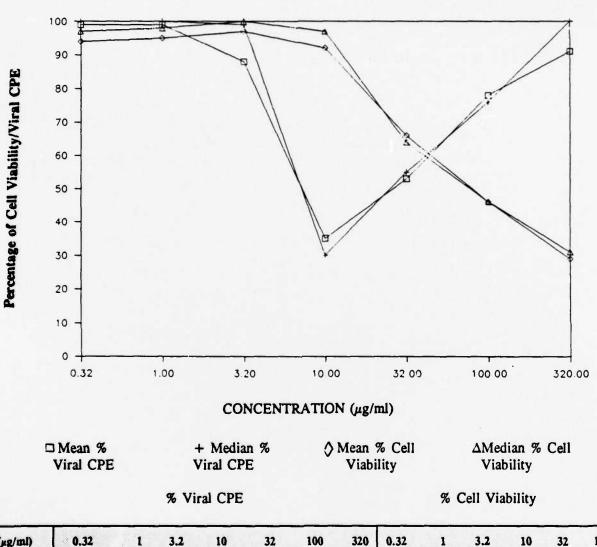
The 42 control tests performed with 2-Thio-6-Azauridine gave a mean Total Antiviral Index (TAI) of 17.10% (SD \pm 12.40) and the median value was 14.30%. The TAI measures the overall antiviral effectiveness of the compound and it ranged from ~ 0 - 50.23% during this period. The mean Selectivity Index (SI) was 5.34 (SD \pm 7.70) and the median SI value was 3.40, indicating moderate antiviral selectivity for 2-Thio-6-Azauridine against SF virus. The SI ranged from ~ 0 - 45.73 during this period. However, the closeness of the mean and median values indicate that the present execution of the SOP is consistent and repeatable.

The mean Antiviral Index 25% (AI₂₅) value was 8.80 (SD \pm 11.70). The median AI₂₅ value was 5.20 (range 0.150 - 71.53). The mean Antiviral Index 50% (AI₅₀) was 11.00 (SD \pm 10.5) with a median of 9.20 (range 0 - 51.58). The mean Antiviral Index 95% (AI₅₀) was 1.10 (SD \pm 3.20), with a median of 0 (range 0 - 11.53). This indicates that the, 2-Thio-6-Azauridine, does not consistently reach 95% antiviral reduction levels.

The mean Antiviral Inhibitory Concentration 25% (IC₂₅) was 4.90 μ g/ml (SD \pm 2.30). The median IC₂₅ value was 4.60 μ g/ml (range = 1.24 - 12.80 μ g/ml). The mean Antiviral Inhibitory Concentration 50% (IC₅₀) was 6.00 μ g/ml (SD \pm 5.40). The median IC₅₀ value was 5.90 μ g/ml (range = 0 - 26.80 μ g/ml). The mean Antiviral Inhibitory Concentration 95% (IC₉₅) was 1.50 μ g/ml (SD \pm 4.80). The median IC₉₅ value was 0 μ g/ml (range from 0 - 26.2 μ g/ml). This discrepancy indicates that the 2-Thio-6-Azauridine does not consistently reach 95% reduction levels. During this reporting period, the highest starting concentration of 2-Thio-6-Azauridine (320 μ g/ml) was varied to properly evaluate the maximum antiviral effect of 2-Thio-6-Azauridine. The best window appears to be from a high concentration of 100 μ g/ml to a low concentration of 0.320 μ g/ml in six 1/2 log₁₀ increments. (See Figure 37-B). At this scale (0.32 - 100 μ g/ml) all important antiviral (IC₂₅, IC₅₀, IC₉₅ and TAI) parameters are measured as well as all important cytotoxicity parameters (TC₂₅, TC₅₀, except TC₉₅) as also indicated.

The average maximum antiviral inhibitory level of 42, 2-Thio-6-Azauridine, tests (Figure 37-B) was reached at 10 μ g/ml of the compound with 70% antiviral effect. Maximum antiviral effect (~70%) was found with a simultaneous ~5% cytotoxic suppression. Above 10 μ g/ml concentration, 2-Thio-6-Azauridine starts to lose its antiviral potency (~10%) at 320 μ g/ml, while simultaneously the 2-Thio-6-Azauridine becomes maximally toxic (~70%) with increasing cytotoxicity.

2-THIO-6-AZAURIDINE - VS- SF VIRUS



Conc.(µg/ml)	0.32	1	3.2	10	32	100	320	0.32	1	3.2	10	32	100	320
Mean	99	99	88	35	53	78	91	94	95	97	92	66	46	29
Median	100	100	99	30	55	76	100	97	98	100	97	64	46	31
Std. Dev.	0.02	0.03	0.2	0.22	0.23	0.14	0.11	0.07	0.06	0.05	0.11	0.16	0.14	0.08

Figure 37-B
Average Antiviral and Cytotoxicity Values for 42 Positive Control Compound Tests

4.1.7.1.2 <u>Maximum Antiviral Effect of 2-Thio-6-Azauridine vs SF Virus:</u> Since the metabolic activity of the cells was an unknown function during the testing period, it was monitored indirectly by measuring the maximum antiviral effect of the control compound 2-Thio-6-Azauridine. This demonstrated the amount of infectious virus produced by the cells (Maximum Percent CPE).

A bar graph scatter plot (Figure 38-B) depicts the distribution of the maximum antiviral reduction values of all 42 control compound assays for 2-Thio-6-Azauridine. The results indicate that the average maximum antiviral reduction obtained with the present SOP is around 70% (SD \pm 18.70) reduction levels. The maximum reduction levels vary from 40 - 100% but remain quite consistently around the median of 72%. The assay control values give a shifted half-bell-shaped distribution curve toward the maximum 100% reduction level. This indicates quite a consistent day-to-day performance of the control compound in the SF-MTT assay.

Recommendations:

Based upon the data obtained in parallel studies with Ribavirin, we recommend that 2-Thio-6-Azauridine (AVS #6724) will be used as a second control compound against SF virus. It's overall performance is slightly poorer than the present control, Ribavirin. However, it is readily available from Sigma Chemical Company, it is inexpensive and works at ~5 - 10-fold lower concentrations than Ribavirin.

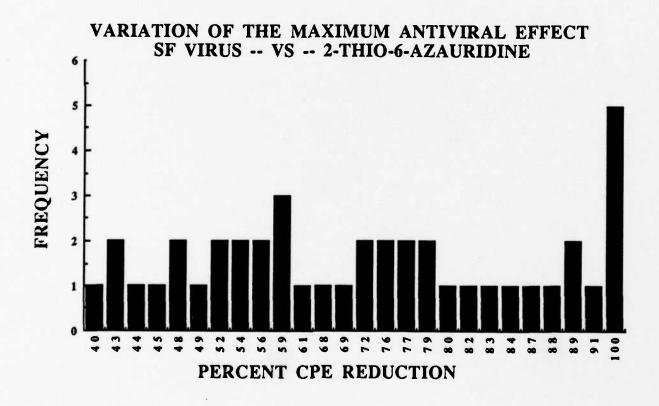


Figure 38-B

Maximum Antiviral CPE Reduction (%).

Summary of 42 Control Tests.

4.1.7.1.3 Cellular Cytotoxicity of 2-Thio-6-Azauridine vs SF Virus:

SF-Control Compound-Cytotoxicity Performance: The 42 cytotoxicity values of the positive control compound 2-Thio-6-Azauridine are also very consistent. The mean cell Toxic Concentration 25% (TC₂₅) was 31.10 μ g/ml (SD \pm 53.30) and the median was 25.5 μ g/ml (range of 0.81 - > 100 μ g/ml). The mean cell Toxic Concentration 50% (TC₅₀) value was 73.80 μ g/ml (SD \pm 27.40) and the median was 84.60 μ g/ml (range of 21 - > 100 μ g/ml). The mean cell Toxic Concentration 95% (TC₉₅) value was 147.10 (SD \pm 91.40) and the median was 100 (range of 100 - 320 μ g/ml). This discrepancy indicates that the control compound 2-Thio-6-Azauridine does not consistently reach 95% cytotoxicity levels.

As can be seen from Figure 37-B, the toxicity starts to become measurable above the concentration of 10 μ g/ml and the maximum toxicity has not been reached at 320 μ g/ml. Further increase of the concentration of 2-Thio-6-Azauridine would be needed to properly evaluate the maximum cytotoxicity of 2-Thio-6-Azauridine.

Also, Figure 37-B indicates that when the cytotoxicity reaches ~ 10% at 10 μ g/ml, the control compound (2-Thio-6-Azauridine) has reached simultaneously its maximum antiviral effect (70%). The cytotoxic effect of 2-Thio-6-Azauridine is insignificant between 1 and 10 μ g/ml. The average cytotoxicity reached ~ 70% at 320 μ g/ml, which is the highest 2-Thio-6-Azauridine concentration tested.

2-Thio-6-Azauridine has a definite cytotoxic suppression on cellular metabolism and growth. However, the TC_{25} and TC_{50} toxicity could be consistently achieved with the 100 μ g/ml concentration of 2-Thio-6-Azauridine. Therefore, a readjustment to 320 μ g/ml as being the highest 2-Thio-6-Azauridine concentration tested is not needed. However, at this concentration (320 μ g/ml) the TC_{95} cannot yet be measured consistently.

4.1.7.1.4 SF-Assay Plate Quality Controls: Cell Load and Virus Load Parameters 2-Thio-6-Azauridine: The MTT assay is fundamentally dependent upon the quality of the assay plates. Our large-scale antiviral testing is dependent upon the uniformity of the test plates produced for the daily assays. Equal loads of cell load and virus load as well as the consistent performance of the reagents used daily was monitored. A sample of the plate variation control for the period of November, 1989 through January, 1991 is presented in Figures 39-B, 40-B, and 41-B.

SF-Control Compound-Cell Load Performance: A bar graph scatter plot of the mean cell control (O.D. reading) of 42 control assays is plotted in Figure 39-B. The results indicate that the cell O.D. readings reached a mean of 0.990 (SD \pm 0.160) with a median of 1.000 (range of 0.420 - 1.240). This indicates that a uniform and equal number (18,000 cells/well) of cells are being loaded into every well in the 96-well plate during the day-to-day operation. The cells reduced MTT to formazan giving maximum blue color uniformly and consistently.

SF-Control Compound-Virus Load Performance: A bar graph scatter plot of the mean virus load O.D. readings of the 42 control assays is presented in Figure 40-B. The results indicate that the average virus load O.D. reading reached a mean of 0.300 (SD \pm 0.120 with a median of 0.300 (range of 0.090 - 0.650). This demonstrates that a good cell destruction is taking place and a uniform load of virus (32 TCID₅₀) is administered on the cell monolayer with very consistent viral CPE results.

SF-Control Compound-Assay Differential Performance: A bar graph scatter plot of the mean O.D. differential values of the 42 control assays is provided in Figure 41-B. The results indicate that the average differential O.D. reading is 0.690 (SD \pm 0.170) with a median of 0.720 (range 0.103 - 0.965). The single bell-shaped curve is reasonably sharp and uniform. This reflects that the assays are executed consistently and are repeatable during day-to-day operation with close to 72% measurement accuracy.

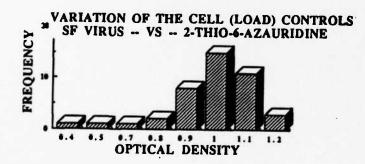


Figure 39-B

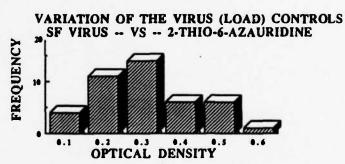


Figure 40-B

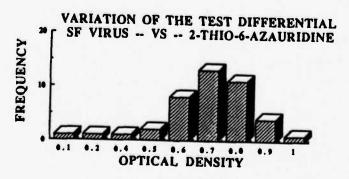


Figure 41-B

4.1.7.2 SF-Antiviral Activity Results:

New Drugs with 95% Antiviral Reduction Levels: Out of the 6487 actual single drug tests, 95 new compounds demonstrated excellent antiviral activity, having antiviral reduction values of equal to or better than 95%. This represents around 1.5% of the test compounds being active at this excellent reduction level. These compounds are summarized in Table 19 according to the highest Total Antiviral Index (TAI). Compound AVS-5580 demonstrated the greatest in vitro promise, having a TAI of 97% and Selectivity Index (SI) of >313. The next 22 compounds demonstrated excellent antiviral activity with TAI's greater than 50% and SI values that ranged from <1 - >320. Forty other compounds demonstrated good antiviral activity, having TAI's that range from 25 - 48% and SI's from <1 - 18. The rest (32) compounds had only moderate activity with TAI's ranging from 5 - 24 and SI's that ranged from 0.03 to 5.2.

It is worthwhile to note that compounds received in shipment number 62 were mostly colored (Table 19). Therefore those compounds appearing in the 95% active category from shipment number 62 should be interpreted with caution, since colored compounds create false positive readings with the MTT assay.

Table 19 ${\tt AVS~Compounds~Active~Against~Sandfly~Virus~(SF)~at~AI_{95}~Level}$

				225									
		AVS	Ship-		Diff-			0-					
Vlr	us	No.	ment#	Date	rnt1.	IC 95		TC 95		AI 95	SI		TAI
	CB	EEOO	54	05/02/00	1 120	700 00 5		100000		107.00 -	212 00		07 00
		5580 5278	52	05/02/89		789.00		320.00		127.00 >			
		5278	52 52	03/21/89		23.90				13.40 >			
				03/21/89		8.55 >				37.40 >			
		1644	64	05/03/90				1000.00		35.78	68.37		
		4452	44	08/16/89	0.609	21.10		320.00		15.15			
		4871	61	11/29/89				1000.00		34.04			
		6417	66	07/06/90			>	1000.00	>	34.38	49.71	>	
		5241	52	12/14/89		28.70		966.00		33.69	45.88		64.58
		4796	46	01/12/89		8.10 >		320.00		39.50	94.70		
		5250	52	03/14/89		86.70					320.00		
		4855	61	11/29/89		29.60				10.80 >			
	_	5643	57	08/16/89	0.690	31.40 >				10.20 >			
		4611	65	07/06/90		277.00				3.61	62.74		
		2320	53	04/11/89				320.00		23.00	49.00		
		6714	67	07/12/90		409.00 >				2.44 >			
	7.1	4763	44	11/02/88		68.40 >		320.00		4.68			52.59
		5040		02/08/90		280.00				11.41	28.63		
		0217	33	04/19/89		79.60 >		320.00		4.02 >			
		4822	48	12/14/89		8.03 >		320.00		39.84	25.63		
		4465	45	12/06/88		192.00 >				1.66 >			
		0206	4	12/05/89		89.20 >				11.21	18.43		
		5067	48	12/14/89	0.559	155.00 >				6.47	24.62		
		5075	48	03/01/89		65.20	>	320.00	>	4.91 >			
		5137	57	08/16/89		44.30		309.00		6.97	13.44		
		5601	62	12/05/89		298.00 >	>		>	3.35	17.74		
		4991	51	03/14/89		31.60		340.00		10.80			46.17
		1217	52	03/15/89		74.10		320.00		4.32	9.12		44.43
		5058	62	12/05/89		289.00 >	> :		>	3.46 >			44.29
		4427	44	03/29/89	0.810	2.91		75.70		26.00			43.67
		6986	68	08/07/90	0.596	88.10		963.00		10.93	15.97		41.73
		4757	44	11/02/88		222.00 >			>	1.44 >			41.63
		5138	57	08/16/89	0.536	93.80		1000.00		10.66			40.15
		7084	72	08/30/90		8.32		96.30		11.57	12.21	>	38.97
	-	5271	52	12/14/89		282.00 >	>		>	3.54	12.65		38.74
		4939	51	03/07/89	0.842	10.00		83.80		8.38 >	10.80	>	37.32
		3802	35	09/12/89	0.786	0.30 >	>	10.00	>	33.49	5.51	>	37.07
		4934	51	03/07/89		2.98		80.00		26.90	15.40	>	36.63
		4438	44	04/05/89		8.22		30.70		3.73			36.13
	SF	7083	72	08/30/90	0.718	31.10		304.00		9.78	10.22		
	SF	4753	44	12/14/89	0.547	2.94		30.50		10.37	6.89		35.50
		8271	76	12/12/90	0.803	280.00		979.00		3.50	6.33	>	35.25
	SF	6315	63	10/04/90		3.16		30.90		9.78	8.79	>	34.13
	SF	4071	62	12/05/89	0.622	302.00 >	>	1000.00	>	3.31	10.47	>	33.61
		6977	68	08/02/90	0.736	89.00		309.00		3.47			32.48
	SF	6788	67	07/12/90	0.800	947.00	>	1000.00	>	1.06	0.27	>	32.16
	SF	7092	72	10/25/90	0.569	1070.00	>	1000.00	>	0.94	0.03	>	32.11
	SF	3688	32	11/10/88	1.011	95.40 >	>	320.00	>	3.36			31.82
	SF	6209	62	01/16/90		291.00 >	>	320.00	>	1.10 >			31.54
	SF	7085	72	08/30/90		9.06		100.00		11.03	6.44		30.82
	SF	4747	44	11/01/88		27.50		96.60		3.52	7.16		30.80
		5532	56	08/09/89		293.00		966.00		3.30			30.01
		2275	53	04/11/89		86.70 >	>	320.00	>	3.69	6.75		29.77
		6477	66	07/10/90		28.30		288.00		10.15			28.71
		6482	66	07/10/90		28.70		97.30		3.39			28.03
		1645	33	11/01/90		29.60 >	>	100.00	>	3.38			27.60
		5498	53	04/18/89		287.00 >		320.00		1.11 >			27.25
		5905	61	10/31/89		3.20		30.30		9.47 >			27.06
			_	-,,		J. -							

Table 19 (Cont'd)

	AVS	Ship-	Test	Diff-				
Virus	No.	ment#	Date	rntl.	IC 95	TC 95	AI 95	SI TAI
SF	4990	51	03/14/89	0.155 <	1.00	198.00 >	198.00 >	4.52 > 26.19
SF	4240	39	09/12/89	0.614	9.70	30.90	3.18	5.69 25.90
SF	6194	62	01/31/90		290.00 >	320.00 >	1.10 >	2.64 > 25.76
SF	6184	62	01/09/90		283.00 >	320.00 >	1.13 >	3.51 > 25.70
SF	6179	62	01/09/90		28.70	96.60	3.36	4.50 25.54
	6202	62	01/31/90		294.00 >	320.00 >	1.09 >	2.35 > 24.85
	2453	64	03/13/90		9.18	30.90	3.36	3.63 > 23.92
SF	6837	68	08/02/90		89.60	306.00	3.41	3.16 23.76
	7087	72	10/02/90		9.30 >		10.75	4.04 23.48
	5253	52	03/15/89		2.62	29.80	11.40 >	4.81 > 23.45
	5495	53	04/18/89		2.83 >		113.00 >	5.16 > 23.12
	6225	62	11/01/90		295.00	966.00	3.27	3.40 > 23.09
_	6195	62	12/05/90		287.00	963.00	3.35	4.09 22.90
	7086	72	08/30/90		2.91	29.90	10.26	4.94 > 21.69
- 9	6214	62	01/16/90		302.00 >		1.06 >	1.79 > 20.87
	8370	76	12/13/90		320.00	2260.00	7.05	4.06 20.33
	6219	62	11/01/90	0.791	286.00 >		1.12 >	3.06 > 20.26
	6217	62	01/16/90		302.00 >		1.06 >	1.79 > 20.06
	6976	68	09/13/90		90.60	308.00	3.39	2.48 19.75
	5531	56	08/09/89		96.00	320.00	3.33	3.05 > 19.69
-	6979	68	08/07/90		315.00	966.00	3.07	3.16 19.32
	7032	69	07/31/90		290.00 >		1.10 >	2.68 > 18.27
	7457	73	09/20/90		3.02	27.90	9.24	3.46 > 17.22
	6197	62	01/31/90		301.00 >		1.06 >	1.86 > 16.97
	4754	44	11/02/88		2.95	9.66	3.27	3.43 > 16.63
	7042	69	07/31/90		94.20	309.00	3.28	2.82 15.92
	7049	69	08/02/90		296.00 >		1.08 >	2.19 > 15.67
	7073	72	10/25/90		312.00	986.00	3.16	2.66 > 15.48
	6207	62	10/30/90		302.00	966.00	3.20	2.74 15.08
	7458 6198	73 62	09/20/90	0.706	2.96	9.61	3.24	2.82 > 14.86
		62	01/31/90		300.00 >	320.00 > 320.00 >	1.07 > 1.06 >	1.91 > 14.47 1.79 > 13.78
	6201	_	01/31/90		302.00 >			1.94 > 11.23
	7430 7003	70 69	08/29/90		300.00 >		1.07 >	1.86 > 11.23
	5242	52	07/26/90		301.00 >		1.06 >	1.91 > 10.88
	7418	70	03/08/89 08/23/90		300.00 > 302.00 >		1.07 > 1.06 >	1.81 > 10.88
	4074	48			_			1.89 > 6.22
		69	12/07/89			26.50 >	26.50 >	
SF	7023	9	07/31/90	0.749	302.00	88.30	0.29	0.03 > 5.04

New Drugs with 50% Antiviral Reduction Levels: Out of the 6487 actual single drug tests, 283 new compounds demonstrated good antiviral activity, having antiviral reduction values equal to or better than 50%. This represents around 4.4% of the test compounds being active at this good antiviral reduction level. These compounds are summarized in Table 20 according to the highest Total Antiviral Index (TAI). AVS-5277, 2951 and 4098 demonstrated the best TAI of 63, 52 and 47% and SI is of > 150. Twenty other compounds demonstrated moderate antiviral activity, having TAI's that ranged from 31 - 43% and SI's from 3 - 173. The rest (260 compounds) showed marginal antiviral activity with TAI's that ranged from <1 to 29% and SI's from <1 - > 320.

It is worthwhile to note (Table 20) that compounds received in shipment number 62 were mostly colored. Therefore those compounds appearing in the 50% active category from shipment number 62 should be interpreted with caution, since colored compounds create false positive readings with the MTT assay.

Table 20 $$\operatorname{AVS}$$ Compounds Active Against Sandfly Virus (SF) at AI_{50} Level

1	AVS	Ship-		Diff-	70.50			
Virus	No.	ment#	Date	rntl.	IC 50	TC 50	AI 50	SI TAI
SF	5277	52	12/14/89		6.26 >	1000.00 >	159.87 >	159.87 > 63.70
SF	2951	26	09/07/89	0.658	0.21 >	32.00 >	151.74 >	151.74 > 51.85
SF	4098	37	02/24/89	1.041	0.00 >		201.00 >	201.00 > 47.49
SF	1019	28	03/06/90	0.575	2.24 >	100.00 >	44.59	26.67 > 43.49
SF	5794	59	10/10/89	0.542	34.50 >		9.28 >	9.28 > 42.19
SF	3592	ABEBE	01/11/89	0.509	26.10 >	320.00 >	12.20 >	12.20 > 41.57
SF	2948	26	09/07/89	0.658	1.60 >	320.00 >	199.84	173.25 > 40.53
SF	1850	53	04/11/89	0.666	10.40 >	100.00 >	9.61	7.42 > 38.87
SF	0919	52	03/15/89		2.30	162.00	70.60	35.90 38.47
	4989	51	03/14/89	0.182	2.28 >			141.00 > 36.16
	5834	59	01/24/90		201.00 >		1.59 >	1.59 > 36.01
SF	5793	59	10/10/89		58.50 >		5.47 >	
	4472	45	10/11/89		57.70 >		5.54 >	
	3621	32	11/09/88		42.50 >		7.52	5.91 > 33.34
	4875	46	01/18/89		64.90 >		4.93 >	
	2980	25	09/07/89		0.20	3.09	15.60	8.83 > 32.54
	2433	17	08/30/89		10.50 >		30.55	9.55 > 31.95
	4984	51	03/14/89		7.93	228.00	28.70	14.50 > 31.50
	3615	32	11/08/88		56.60	262.00	4.63	3.20 > 31.48
	8353	75	11/29/90		33.30 >		9.60	
	0002	46	01/24/89		51.60 >		6.21 >	
	7382	70	08/21/90			1000.00 >	8.91 >	
	6234	GABSN	,,			3200.00 >	6.61 >	6.61 > 30.51
	4439	44	10/11/89		7.74	65.30	8.44	6.20 > 29.28
	5283	52	03/21/89		2.97	251.00	84.60	38.20 > 29.05
	6205	62	01/31/90		75.20 >		4.25 >	
	4473	45	10/11/89		99.40 >		3.22 >	
	5186	58	10/04/89		49.90	887.00	17.76	8.30 > 27.60
	4764	44	11/08/88		106.00 >		3.02 >	3.02 > 27.25
	5369	54	05/02/89		9.09 >		35.20	18.50 > 27.11
	4432	44	04/05/89	0.580	20.30 >		4.92 >	4.92 > 27.00
	5507	53	04/18/89		42.50	309.00	7.27	3.92 > 26.77
	5515	53	04/18/89		10.80	53.90	4.99	2.38 > 26.74
	8499	75	11/29/90		58.10 >		5.51 >	
	3619	32	11/09/88		93.10 >		3.44 >	
	5625	57	06/28/89		64.80 >		4.94 >	
	3584	32	11/03/88		17.90 >		5.59	4.32 > 25.91
	8348	75	11/29/90		48.40	216.00	4.45	3.00 25.89
	6753	67	07/12/90			1000.00 >	11.60	8.80 25.64
	1159	52 45	08/23/89		32.90 75.20 >	234.00 320.00 >	7.12 4.25 >	5.08 25.57 4.25 > 25.57
	1730	61	11/22/88 02/01/90					
	5998	66	07/06/90		3.69 > 32.00	32.00 > 82.70	8.67 2.59	2.51 > 25.57 1.79 > 25.16
	6412 5072	48	03/01/89		19.50	279.00	14.30	7.21 > 25.16
	4035	65	07/06/90				5.42 >	5.42 > 25.10
	4770	44	11/08/88		27.10	1000.00 >	6.60	4.00 > 24.98
	6236		02/08/90			3200.00 >	5.85 >	5.85 > 24.69
	0148	2	08/23/89		0.32 >		10.00	6.84 > 24.56
	2274	12	08/30/89		18.80 >	_	5.31 >	5.31 > 24.52
	5041		02/08/90			1000.00 >	4.79 >	4.79 > 24.40
	5450	53	04/12/89		2.81	29.30	10.40	6.33 > 24.40
	5534	56	06/14/89		62.80 >		5.10 >	5.10 > 24.30
	6218	62	12/05/90		20.50 >		15.58	7.93 24.28
	6373	63	02/27/90		78.50 >		4.07	2.87 > 24.26
	6321	63	03/15/90		2.81	21.10	7.51	5.53 > 23.75
	2363	15	08/30/89		7.79	63.10	8.09	5.40 > 23.58
	2151	52	03/15/89		48.40 >		6.61	3.54 23.52
31	5191	32	03/13/03	5.565	40.40 >	320.00 >	0.01	23.32

Table 20 (Cont'd)

	AVS	Ship-	Test	Diff-								
Virus	No.	ment#	Date	rntl.		IC 50		TC 50		AI 50	SI	TAI
SF	7469	73	10/30/90	0.815		90.20		777.00		8.62	5.53	23.47
	7910	75	11/06/90				>	3200.00	>	5.89	5.33	> 23.08
	4241	46	01/24/89			2.48		22.90		9.26		> 23.02
	3558	31	03/29/89	0.732		30.00		182.00		6.07	3.39	22.91
	5497	53	04/18/89	1.086		88.40	>	320.00	>	3.62 >		> 22.90
	4943	51	03/07/89			1.71		22.50		13.20		> 22.87
	8251	76	12/11/90			50.80		618.00		12.16		> 22.87
	1992	56	06/13/89			94.70	>	320.00	>	3.38 >		> 22.74
	6176	62	01/09/90			75.20		320.00		4.25 >		> 22.63
	6191	62	01/31/90	0.689		96.10		320.00		3.33 >		> 22.58
	4769	44	11/08/88	0.914		14.40		92.40		6.41		> 22.35
	3690	32	11/10/88			32.00	>	100.00	>	3.12 >		> 22.15
	5372	54	05/02/89			68.40		180.00		2.63		> 21.97
	2315	13	12/21/88	0.924		100.00	>	100.00	>	1.00		> 21.92
	3803	35	12/13/88		<	1.00		19.20		19.20 >		> 21.91
	5070	48	03/01/89			1.00		142.00		142.00 >		> 21.74
	5508	53	04/18/89		-	45.70		184.00	•	4.02		> 21.73
	8347	75	11/29/90			273.00	>	320.00	>	1.17 >		> 21.61
1 - 1 - 1 - 1	2631	19	09/06/89			25.60		320.00		12.51		> 21.49
	5818	59	01/31/90			49.60	-	237.00	•	4.78		> 21.49
	7332	70	09/13/90			42.10		204.00		4.85		> 21.06
	3689	32	11/10/88			22.20	>	100.00	>	4.51		> 20.88
	5494	53	04/18/89			19.60		100.00		5.09		> 20.88
	6943	69	09/06/90			49.70		541.00		10.88	6.27	20.74
	7459	73	09/20/90			2.35		18.70		7.98		> 20.72
	7484	73	10/30/90			76.30		849.00		11.12		> 20.60
	5525	56	08/08/89	1.319		52.10		424.00		8.13	4.03	20.56
	5791	59	10/10/89			32.00		204.00		6.38		> 20.48
	7439	73	10/25/90			70.80		646.00		9.12		> 20.27
	4744	44	11/01/88			9.07		63.60		7.02	4.84	20.22
T	4617		02/08/90				>	3200.00	>	5.66	4.04	20.12
	4988	51	03/14/89		<	1.00		184.00		184.00 >		
SF	4785	46	01/12/89			1.00		25.80	>	25.80 >	7.20	> 20.09
SF	3935	35	12/13/88			19.90		175.00		8.80	4.81	19.88
SF	4993	51	03/14/89			3.09		202.00		65.30	36.60	> 19.88
SF	5548	56	08/09/89			16.90		173.00		10.27	5.92	> 19.85
SF	6946	69	09/06/90	0.839		137.00		558.00		4.08	2.46	19.85
SF	4768	44	11/08/88			16.80		137.00		8.15	3.43	> 19.77
SF	1976	1	08/30/89	0.723		16.90	>	100.00	>	5.92	4.41	> 19.58
SF	5795	59	10/10/89	0.619		9.11		198.00		21.71	5.58	> 19.57
SF	5780	59	10/10/89			54.90		222.00		4.05	2.94	> 19.44
SF	5790	59	10/10/89	0.570		56.60	>	320.00	>	5.66	1.34	> 19.27
SF	7068	72	08/30/90	0.721		51.90		208.00		4.01	2.92	19.21
SF	4937	51	03/07/89	0.865		16.80		144.00		8.59	3.92	> 19.18
SF	5520	56	06/13/89	1.099		164.00	>	320.00	>	1.95 >		> 19.11
SF	2318	13	08/30/89	0.652		0.50		10.00		20.01	4.02	19.09
SF	5543	56	08/09/89			0.23		1.81		7.73	4.52	18.77
SF	4919	46	02/01/89			33.30		184.00		5.51	3.01	18.70
SF	5678	57	08/16/89	0.699		64.50		404.00		6.25	2.69	18.63
	6200	62	12/05/90	0.583		10.00	>	320.00	>	32.00 >	32.00	18.63
	5782	59	10/10/89	0.547		65.70		216.00		3.28		> 18.50
SF	6570	66	05/30/90			66.40		320.00	>	4.82 >		> 18.41
	3624	32	11/09/88			256.00	>	320.00	>	1.25		> 18.39
	1355	64	05/03/90			170.00		879.00		5.17	3.38	18.36
F 47 T 44 T	5626	57	06/28/89	0.983		61.60		308.00		4.99		> 18.19
	3613	32	11/08/88			41.00	>	320.00	>	7.81		> 18.15
	2811	48	02/07/89			0.06		0.26		4.52		> 18.08
	6073	62	12/21/89			5.18		21.00		4.06		> 18.01
SF	5285	52	03/21/89	0.684		5.24		20.50		3.92	2.83	> 17.97

Table 20 (Cont'd)

	AVS	Ship-	Test	Diff-										
rirus	No.	ment#	Date	rnt1.		IC 50		TC 50		AI 50	SI		TA.	I
SF	3677	32	11/09/88			8.46		10.00	>	1.18	1.18	>	17.80	0
	6988	68	06/21/90			71.40		320.00		4.48			17.8	
-	4459	45	11/23/88			79.60	>	320.00	>	4.02 >		>	17.7	4
	5503	53	04/18/89			22.80		155.00		6.81	3.19		17.5	
SF	8293	75	11/28/90	0.818		157.00		320.00	>	2.03 >	2.03	>	17.4	1
	4435	44	04/05/89	0.590		95.70		100.00		1.04 >			17.4	
	2506	21	09/06/89			2.16		10.00		4.64			17.0	
	4981	51	03/14/89			1.85	>	320.00	>	173.00 >				
	8374	76	01/16/91			554.00		2050.00		3.71			16.8	
	4995	51	03/14/89	0.232		7.96		52.90		6.65		>	16.7	
_	5142	57	08/16/89			61.10		210.00		3.43	2.54		16.6	_
	5500	53	04/18/89			1.25		5.45		4.36		>	16.4	
	5339	53	04/12/89			26.50		188.00		7.10	3.48		16.29	
	8358	76	01/18/91			54.10		239.00		4.42	3.13		16.20	
	7434	70	10/02/90			174.00		580.00		3.34	2.13		16.2	
	5284	52	03/21/89			182.00	>	320.00	>	1.76 >		>	16.2	
	7935	75	11/08/90			176.00		762.00		4.34	3.08		16.2	
	3684	32	11/10/88	1.011		100.00		320.00		3.20 >			16.1	
	9121	77	01/31/91	0.636	1			3200.00		1.72 >			16.1	
	5367	54	05/02/89			81.10	>	320.00	>	3.95		>	16.0	
	4739	44	07/06/89			15.10		62.60		4.15	2.88		16.0	
	5879	61	10/31/89			27.50		171.00		6.24	3.41		15.99	
_	2503	21	09/06/89	1.121		0.59		6.60		11.13	3.43		15.8	-
	6130	62 48	12/28/89			17.60		68.70		3.90 6.63		>	15.68	
	0360 5652		02/07/89			0.09		0.61			2.74		15.44	-
	0345	57 2	07/06/89 05/10/89			30.90		129.00		4.19 5.33			15.3	
	2563	48	02/07/89			5.02		3.20		0.64			15.19	
	2979	48	02/07/89			1.79		4.48		2.50			15.12	
	6387	69	07/17/90			19.20		66.00		3.45	2.56		14.9	_
	7461	73	09/20/90	0.623		17.90		83.50		4.67	3.23		14.9	
T 1/2 2 2	7048	69	08/02/90	0.776		57.40		203.00		3.54		`	14.79	
	7378	70	08/21/90			194.00		660.00		3.40	2.52		14.7	
	6945	69	09/06/90			95.00		212.00		2.24	1.64		14.4	
	8378	76	12/13/90			170.00		618.00		3.64	2.52		14.3	_
	5350	53	04/12/89			17.50		61.70		3.52		>	14.28	
	8311	76	12/12/90	0.734		224.00		962.00		4.29			14.20	
	6707	67	07/12/90			7.30		76.60		10.50	6.68		14.09	
	5274	52	03/21/89			4.86		13.30		2.74		>	13.9	
SF	5252	52	03/14/89		<	1.00		25.20	>	25.20 >	13.10			
SF	6226	62	01/16/90			231.00	>	320.00	>	1.38 >			13.83	
SF	5528	56	06/13/89	1.294		6.37		21.70		3.41	2.49		13.69	
SF	4592	61	11/29/89			21.50		67.50		3.15	1.41	>	13.40)
SF	5542	56	06/14/89			1.83		7.06		3.86	2.81	>	13.36	5
SF	6435	66	07/06/90	0.771		20.30		92.40		4.55	0.03		13.26	5
SF	1980	27	05/16/89			24.60	>	100.00	>	4.06	2.29	>	13.0	3
SF	3038	28	09/07/89			0.06	>	3.20	>	50.48	4.08		12.88	3
SF	4765	44	11/08/88			89.20		181.00		2.03	0.10		12.82	2
	6944	69	09/06/90			61.60		223.00		3.62	1.56	>	12.83	1
	6212	62	01/16/90			158.00		320.00		2.02 >	2.02	>	12.74	4
	2812	48	02/07/89			0.01	>	0.01	>	1.11 >		>	12.6	
	6029	61	02/01/90			65.70		209.00		3.18	2.33		12.59	
	7072	72	10/25/90			239.00		630.00		2.63			12.5	
	5485	66	05/08/90			179.00	>	320.00	>	1.79 >		>	12.47	
	4436	44	04/05/89			24.40		54.00		2.21	1.20		12.40	
	6223	62	01/16/90			264.00	>	320.00	>	1.21 >			12.34	
	5280	52	03/21/89			1.79		5.21		2.91	1.58			
	7424	70	08/23/90			162.00	>	320.00	>	1.97 >	1.97			
SF	8586	76	01/23/91	0.817		44.30		256.00		5.78	2.52	>	12.19	•

Table 20 (Cont'd)

	AVS	Ship-		Diff-				
Virus	No.	ment#	Date	rnt1.	IC 50	TC 50	AI 50	SI TAI
SF	7333	70	09/13/90	0.732	67.60	204.00	3.01	2.05 > 12.03
	2449	18	09/06/89		21.10	93.60	4.43	0.47 > 11.99
	8350	75	11/29/90		9.52	21.00	2.21	1.27 > 11.99
100000	3450	32	04/04/89		10.00	19.50	1.95	1.02 > 11.92
	5403	54	05/04/89		28.40	91.60	3.23	2.18 11.87
	4750	44	11/01/88		13.40	44.30	3.31	1.75 > 11.66
	4996	51	03/14/89					320.00 > 11.56
	7373	70	08/21/90		174.00 >		1.84	1.69 > 11.52
	6978	68	06/21/90		187.00 >		1.71 >	1.71 > 11.45
SF	5997	61	11/07/89		2.88	28.30	9.84	2.40 > 11.40
SF	5936	60	10/24/89	0.933	21.40	57.40	2.68	1.62 > 11.35
SF	7365	70	08/16/90	0.660	769.00 >	1000.00 >	1.30	1.00 > 11.31
	5156	57	08/15/89	0.581	1.00	2.68	2.68	1.23 11.24
SF	4751	44	11/01/88	0.971	22.50	56.40	2.51	1.54 > 11.00
	5484	53	12/14/89		198.00	618.00	3.13	2.16 > 10.98
	5653	57	07/06/89		9.39	26.50	2.82	1.60 > 10.87
	5693	57	07/18/89		141.00 >		2.27	1.94 > 10.84
	2586	19	09/06/89		69.50 >		4.60	0.10 > 10.81
	8699	76	01/22/91		179.00 >		1.79	1.41 > 10.70
	4982	51	03/14/89			278.00 >		151.00 > 10.53
175.00	5496	53	04/18/89		80.00 >		1.25 >	1.25 > 10.35
	4051	37	10/04/90		0.29	0.66	2.24	1.64 > 10.28
	5538	56	06/14/89		213.00 >		1.51 >	1.51 > 10.28
	3686	32	11/10/88		50.50	179.00	3.55	2.15 10.26
	4827	48	02/14/89		241.00 >		1.33 >	1.33 > 10.23
	8352 6422	75 66	11/29/90			5.61 >	5.61 >	2.67 > 10.22
	8250	76	07/06/90 12/11/90		211.00 75.70	585.00 205.00	2.78 2.72	1.79 10.19 1.96 > 10.17
	7059	72	10/25/90		184.00	602.00	3.28	1.85 10.15
	4829	48	02/14/89	1 222	252.00 >		1.27 >	1.27 > 10.11
	4756	44	11/02/88		7.52	19.60	2.61	1.79 > 10.00
	4428	44	03/29/89		27.90	63.10	2.26	1.17 9.97
	7487	73	10/30/90		214.00	720.00	3.36	2.43 9.93
	8511	76	12/19/90			3200.00 >	2.30	2.30 > 9.84
	8698	76	01/22/91	0.645	8.75	27.70	3.17	2.16 9.76
SF	4887	46	01/18/89		2.85	15.10	5.29	2.43 > 9.61
SF	5714	58	10/04/90	0.762	2.39	6.05	2.53	1.70 9.61
SF	5906	61	10/31/89			10.40 >	10.45 >	2.72 > 9.54
SF	7356	70	08/16/90		228.00 >	320.00 >	1.41 >	1.41 > 9.43
SF	4800	46	01/17/89	1.452	188.00 >		1.70 >	1.70 > 9.35
	4036	65	04/24/90		223.00 >		1.43 >	1.43 > 9.14
	5307	52	03/22/89	1.044	20.10	60.60	3.02	1.50 8.89
	0147	64	03/13/90		27.40	90.30	3.29	2.18 > 8.50
	4527	47	02/01/89		8.27	17.60	2.12	0.87 > 8.48
	6413	66	05/10/90		23.90	51.00	2.13	0.77 > 8.26
	5247	52	03/14/89			196.00 >	196.00 >	88.50 > 8.04
	6970	68	08/02/90		69.80	186.00	2.66	1.70 > 8.00
	8285	75	11/28/90		147.00 >		2.17	0.47 > 7.83
	8677	76	01/22/91		250.00 >		1.28 >	1.28 > 7.80
	4752	44	11/01/88		5.31	8.68	1.64	1.12 7.74
	5197	58	10/04/90		23.70	56.40	2.38	1.46 > 7.73
	5994	61	11/07/89		256.00 >		1.25	1.00 > 7.66
	6896 7320	69	09/05/90		2.61	6.82	2.61	1.79 7.52
	3491	70 65	08/14/90		78.90 289.00 >	207.00	2.62	1.88 7.45
	4911	46	04/17/90 01/31/89		289.00 > 171.00	320.00 > 284.00	1.11 > 1.66	1.11 > 7.24 0.85 > 7.10
	7055	72	08/29/90		89.60	207.00	2.31	1.67 7.05
	5134	57	08/15/89		23.30	59.90	2.57	1.69 7.04
	7344	70	08/14/90			66.70	3.23	
31	7377	70	20/14/30	0.710	20.60	00.70	3.23	1.55 6.87

Table 20 (Cont'd)

						•						
	AVS	Ship-	Test	Diff-								
Virus	No.	ment#	Date	rntl.		IC 50		TC 50		AI 50	SI	TAI
SF	4951	51	03/07/89	0.549		32.00		72.20		2.26	1.45	6.82
	5539	56	08/09/89			2.59		6.96		2.69	1.71	6.79
	8313	76	12/12/90				>	3200.00	>	1.12 >	1.12 >	6.73
	7323	70	08/14/90					1000.00		1.09 >	1.09 >	6.58
	4434	44	04/05/89			28.20		53.00		1.88	1.00	6.30
	5133	56	08/15/89			7.62		8.91		1.17	0.72	6.30
	8376	76	12/13/90			27.70		89.90		3.25	1.94	6.13
	7057	72	10/25/90			85.00		215.00		2.53	1.51 >	6.09
	6900	69	07/19/90			9.19		19.90		2.17	1.51	5.85
	2589	65	04/12/90			273.00				1.17 >	1.17 >	5.74
	5363	54	06/28/89			10.00		100.00		10.00	0.40 >	5.71
	6675	64				23.40		52.10		2.23		5.69
	2590	19	04/05/90				_				1.24	
		-	09/06/89			191.00		320.00		1.68	0.02 >	5.67
	1337	33	08/29/89			23.50	>	32.00	>	1.36 >	1.36 >	5.53
	5174	58	07/25/89			23.60		64.90		2.74	0.04	5.46
	7427	70	08/29/90		_	249.00	>	320.00		1.28 >	1.28 >	5.46
	5251	52	03/14/89		<			55.50	>	55.50 >	32.70 >	5.34
	7044	69	08/02/90			95.60		194.00		2.02	1.36	5.22
	4762	44	11/02/88			28.50		210.00		7.37	0.09	5.20
-	4746	44	11/01/88			82.70		140.00		1.69	0.23	5.17
7.50	5113	56	08/15/89	0.809		7.57		18.30		2.41	1.50	4.83
	5449	53	04/12/89			10.00		20.40		2.04	0.61	4.73
	7429	70	08/29/90			246.00	>	320.00	>	1.30	0.80 >	4.69
	4859	48	02/21/89			81.40		167.00		2.05	1.08 >	4.67
	5098	56	08/08/89			227.00		443.00		1.95	1.02	4.57
	2600	65	04/12/90					1000.00		1.18	0.34 >	4.33
	7071	72	08/30/90			100.00	>	100.00	>	1.00 >	1.00 >	4.23
	3559	31	03/29/89			95.70		165.00		1.72	0.91	4.04
	5383	54	05/02/89			79.00		169.00		2.14	1.18	4.01
	8541	76	01/03/91			23.60		2.69		0.11	0.04	3.78
	2591	65	04/12/90	0.927			>	1000.00	>	2.31	0.06 >	3.70
	7022	69	07/26/90	0.572		80.20		177.00		2.21	1.32	3.69
	7040	69	09/11/90			259.00		452.00		1.74	0.89	3.59
	7369	70	08/16/90			305.00		320.00		1.05 >	1.05 >	3.56
	7074	72	08/30/90			292.00		320.00		1.10 >	1.10 >	3.39
	7488	73	10/30/90					1000.00		1.24 >	1.24 >	3.33
	6865	68	06/19/90		<	1.00	>	320.00	>	320.00 >	0.86 >	3.32
	6886	69	09/05/90	0.811		2.79		5.28		1.89	0.99	3.29
	3801	35	12/13/88	0.725		23.40		2.20		0.09	0.02	3.27
SF	5153	57	07/11/89	1.141		97.80		169.00		1.73	0.97	3.14
SF	7925	75	11/08/90	0.649		2620.00	>	3200.00	>	1.22	0.38 >	3.07
SF	7940	75	11/08/90	0.537		320.00		547.00		1.71	1.00	3.01
SF	5797	59	10/10/89	0.656		312.00	>	320.00	>	1.02	0.07 >	2.85
SF	5384	54	05/02/89	1.129		27.70		50.70		1.83	0.97	2.73
SF	4891	46	01/18/89	1.265	<	1.00		2.21	>	2.21 >	1.29 >	2.72
SF	2572	65	04/12/90			306.00	>	320.00		1.05	1.02 >	2.47
	5210	58	08/01/89	1.262		27.00		54.20		2.01	0.72	2.46
SF	6906	69	09/05/90	0.916		9.70		18.10		1.87	0.78	2.44
	5774	59	10/10/89			100.00		30.30		0.30	0.20 >	2.30
SF	8332	76	12/13/90			286.00		525.00		1.84	0.66	2.17
	3800	35	12/13/88		<	1.00		2.61	>	2.61 >	1.39 >	1.98
	8326	76	12/13/90			313.00		537.00		1.72	0.91	1.96
	6942	69	09/06/90	0.886			>	1000.00	>	1.13	0.83 >	1.89
	6220	62	01/16/90			313.00		320.00		1.02	0.84 >	1.83
	7110	70	09/13/90			88.50		153.00		1.73	0.75	1.66
	7036	69	09/11/90	1.112		312.00		460.00		1.48	0.70 >	1.25
	4032	65	07/03/90			291.00		443.00		1.52	0.34	0.84
	2902	26	09/07/89			85.50		89.30		1.04	0.70 >	0.60
	4609	48	03/01/89			0.32		0.60		1.87 <	1.00 >	0.34
	,		-0,01,03	51,05		3.32		5.00		2.0/	2.00 >	J.JT

New Drugs with 25% Antiviral Reduction Levels: Of the 6487 actual single drug tests, 492 new compounds demonstrated moderate antiviral activity, having antiviral reduction values equal to or better than 25%. This represents around 8.0% of the test compounds being active at this marginal antiviral reduction level.

In general, when compared to the 95% and 50% antiviral activity categories, the compounds in this (25%) category do not appear to have any significant antiviral promise and probably do not need to be presently confirmed.

4.1.7.3. Confirmatory Assays:

Some of the compounds were sent to us in more than one separate drug shipment. These compounds were tested more than once. Data from the confirmatory assays are summarized in Table 21. If a compound showed $\geq 50\%$ reduction in CPE during this contract period, then it was considered a candidate for confirmatory testing. The confirmatory tests are from active compounds picked up by both the VR and MTT assay testing. Out of 323 confirmatory tests, 228 compounds were confirmed active during this reporting period and the remaining 95 compounds gave conflicting results. The criteria for activity is that the confirmatory test has to show $\geq 25\%$ reduction in CPE. Failure to confirm the activity in these compounds was probably due to differences during the assay conditions:

- 1) In confirmatory assays the concentration range is adjusted to a more accurate semilog scale to maximize the TAI window.
- 2) Differences in the "differential" of the two runs can cause the compound to read positive or negative, falsely. The variability in the differential can cause false positive or negative bias to be introduced into the calculations, thus reflecting the variability in the maximum activity of the compound.
- 3) The metabolic rate, cell and virus load/well, age, and passage number of the cells may cause the above observed variability in the confirmatory results.
- 4) Problems associated with stability and storage of the compound (i.e., different lot numbers, solubility, light sensitivity, hygroscopic, etc.).
- 5) Problems associated with technical execution of large numbers of plates by different technicians.

During this reporting period the overall confirmatory rate against SF was 71%. The conflicting results should be retested at a later date based on the availability of the compound.

4.1.7.4 Recommendations of SF-Actives Based Upon the In Vitro Results with MTT Assay (Vero Cells).

Based upon the in vitro results with the MTT assay (Vero cells) we recommend the following:

- 1) Compounds with the highest TAI in the 95% activity category that have confirmed results with the exception of "colored" compounds should receive the highest priority for further profile studies and *in vivo* animal testing.
- 2) Compounds with the highest TAI in the 50% activity category that have confirmed results with the exception of "colored" compounds should receive the next highest priority for further profile studies and *in vivo* animal testing.

Table 21

Confirmatory Assays for Compounds Active Against Sandfly Fever Virus

۲ د	ب د	+	٠	+	٠	+	+	•	•	+	•	+	+	+	+	+	+	+	٠	+	+	+	+	+	+	+	+	+	+	+	+	
	iype	TH	H	CPE	CPE	CPE	CPE	HT	CPE	CPE	HTT	8	8	8	CPE	95	CPE	365	CPE	CPE	TTH	TTM	CPE	CPE	SPE	CPE	HT	H	H	CPE	CPE	
	Ι¥Ι	31.30	0.00	3.40	0.43	2.60	1.40	2.79	3.70	1.95	3.09	5.50	1.00	4.40	1.10	4.70	0.40	1.90	0.20	1.20	8.50	2.64	3.60	4.10	3.90	0.50	24.56	0.03	20.03	3.40	1.35	
	SI	6.21 >	9.0	3.20	0.00	3.30	0,60	0.00	6.10	0.22	0.00	154.00	0.10	246.00	90.0	246.00 >	0.00	1000.00	0.00	0.12	2.18 >	0.00	4.35	12.90	3.40	1.43	6.84 v	0.00	3.76 >	6.40	1.55	
	A1 95	0.00	0.00	0.0	0.00	0.00	9.1	0.00	0.00	4.20	0.00		0.00		0.00		0.00	•	0.0		0.00	0.00	00.00	20.00	00.00	0.00	0.0	0.0	0.0	0.00	1.00	
	70 95	320.00	00.00	100.00	10.00	10.00	32.00 >	100.00	100.00	100.00	10.00	100.001	320.00	320.00	320.00	320.00	320.00	3.20	10.00	3.20	294.00	320.00	^	٨	A	9.				320.00	320.00	
	10 95	0.00			0.00			0.00			0.00	32.00 >	0.00 - 320.00	0.00	0.00 - 320.00	0.00 > 320.00	0.00	0.00	0.00	0.00		0.00				0.00					320.00	
	AI 50	6.21	0.00	3.20	0.00	10.00	5.60	0.00				154.00	0.10	246.00	90.0	246.00	0.00	00.0001	0.00	0.45	3.29	0.00	13.90 -	12.90 -	3.40	1.43	10.00	0.00	10.44	6.40	1	
	10 50	320.00 >	00.000	32.00	10.00	10.00	32.00 >	8.30	100.00	32.00	10.00	100.00	10.00	320.00	1.00	320.00	32.00				90.30	38.50	32.00	3.20	9.1	1.00	3.20 >	1.00	6.22	320.00 >	100.00	
	10 50	> 09.15		10.00	0.00	.98 >	5.70 >	0.00					72.00		17.00			0.001	0.00	0.87	27.40	0.00	2.30	0.20	0.30	0.70	0.32 >	0.00	0.60		64.17	
	AI 25	13.50	0.00	0.00	0.00	32.00	1.00	0.00	00.00	1.00	0.00		0.10		0.03	000.000	0.10	2.10 <	3.10	3.20	3.78	1.61	21.00	9.60	9.60	1.00	13.90	0.70	2.3	00.00	1.43	
	10 25	320.00 >	\$20.00	32.00	9.90	3.20	3.20	4.30			10.00		3.20	,	0.32 -		10.00	99.0	9.90	2.10	59.90	2.89	21.00	2.10	99.0	99.0	2.19	27.0	5.24	320.00		
	10 25	٨	0.00	0.00	0.00	0.10	3.20	0.00	1.00 >	0.32	0.00	0.10 >	32.00 -		10.00 ~	0.32 >	100.001		2.10	99.0	15.90	1.79	1.00	0.32	0.10	99.0	0.16	19.0	0.39	3.20 >		
	Diff.	1.229	0.581	Y.	K	¥	¥	789.0	¥	¥	0.675	×	· Y	V VI	KA KA	K	××	KA	¥	¥	0.582	0.715	¥.	×	¥	¥	0.750	0.918	0.682	K	¥.	
10	÷	OEY	10K	1	i	1	-	2	;	:	880	1	1	:	;	1	:	;	:	1	020	VIB	1	:	:	:	RB3		2A8	1	-	
	Date	01/24/89	12/20/89	10/28/86	12/01/86	10/28/86	05/17/88	03/28/89	10/28/86	05/17/88	08/23/89	10/08/86	07/27/87	10/08/86	07/27/87	10/08/86	08/18/87	10/08/86	08/18/87	05/18/88	03/13/90 UJQ	05/01/90	10/14/86	11/18/86	12/15/86	08/18/87	08/23/89	05/30/90	10/04/90	10/29/86	05/18/88	
chia	ment of	7 9	94	2	~	2	2	53	2	2	7	_	_	_	_	_	_	_	-	_	3	z	2	7	2	2	7	29	2/67	~	5	
41/6	No. ment	0005	2000	0033	0033	9000	9900	900	1200	1700	1200	6200	6200	7800	9008	7600	7600	0139	0139	0139	0147	0147	0148	0148	0148	0148			0148	7610	0197	

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	Type		2	6	HTH	HTT	TTH	HTT			E :		30	9 6	2	T.	Je	9 6	¥ ;		9	200		<u> </u>	S	307	i g	Ē		CP EP	<u>8</u>	9	H	H	TTH	I	H	9 85	SPE	CPE		Ē			
	TAI	3	2.0	2.20-	31.50	4.66	50.77	5.88		9.5	3.5	6.	2 30	-	2.0	97.0	2 200	907 0	9.0	6.0	2.70	1 60	7. 7.	5.5	1.40	0 200	00	1.78		2.50		1.70	15.44	0.80	0.14	5 80	00.0	4.20	4.30	0.80		16.82		43.49	
	s	3	00.76	09.4	6.84 V	15.16 >	18.43 >	0.00		Z4.40 ×	20.00	· 8.	8	2	3.	8	6	2	0.30	v 00.0	6.90	7 20	200	2.70	3.40	02.0	7 20	0.0		18.00	9.0	0.10	2.74	0.00	0.0	1.00 >	0.00	0.03	16.70	6.7 K	5	2.07		26.67 >	
	A1 95	8	2.50	3.20	3.45 >	11.26	11.21	0.00		× 20.4	88	3.	3.20	8	3	0.00	4	2	8 6	3	31.00	5	8 8	3	0.30	000	8	0.0		9.0	0.00	3.10	0.00	0.00	0.00	7.57	00.0	313.00	31.30	1.00	8	8 8		88.	
	1c 95	6	350.00	320.00 >	320.00 >	000.00	000.000	1000.00		320.00 >	320.00	320.00	320 00	22 00	32.00	100.00	100	2 30	3.50	30.00	10.00	5	22.00	36.00	10.00	5	0	32.00		3.20	3.20	1.00 ×	3.20	1.00	9.6	7.57 >	30.80	٨	100.00	1.00	00	308.00		100.00	
	10 95			٨	٨	٨	٨	0.00 > 1		^	90.0	^	100.00	9	9.0	0.00	22 00		9.0	· 6.6	0.32	1 00 \$	8 6	3	32.00	0		0.00		0.35	0.00	0.32 >	0.00	0.00	0.00	1.00	0.00	0.32 >	3.20 >	1.00 \$	6	8 8		9.6	
nt'd)	A1 50		37.00		3	22.08	26.87	0.00		20.42	26.40	3	80		3	0.00	83 00	2 40	0.30	3	83.00	00 5	2	0.50	7.00	200	25.00	0.0		200.00	10.00	11.00	6.63	0.00	0.00	1.17 <	0.00	0.03	16.70	6.73	9	3.0		44.59	
Table 21 (Cont'd)	TC 50		٨	٨	320.00 >	742.00	860.00	305.00		320.00	320.00	320.00	100.00	23 00	32.00	100.00	100 00	200	2.50	0.92	5.00 ~	1 00 1	8 6	۲۰%	20.00	0.0	5 00 2	10.00		\$	2.00 -	1.00	0.61	0.23	0.89	1.17 >	19.80	3.20	10.00	0.32		195.00	;	100.00	
18	10 50		. 00.0	\$ 25.00	40.80 >	29.60	32.00	0.00		× 01.51	4 60.00	· 00.0	53,50	34 04	0.10	0.00	1 20 -	27.0	9.00	9.0	0.06 ~	0 20 4	22.0	0.30	3.00 -	01.0	0 20 =	0.0		0.01 ~	0.20	0.00	60.0	0.0	0.00	1.00	0.00	0.10	09.0	0.40	,	64.20	i	2.24 >	
	A1 25	9	00.00	10.00	10.00	39.97	48.59	1.22	,	30.00	14.41	5.	2.50	1 15	. 43	0.00	5		3.0	8.0	10.00	10.00	3	2	31.00	10	10.00	1.10		32.00	0.30	0.30	2.40	0.00	29.0	1.00 <	0.00	21.00 <	31.40	0.70		2.9		39.94	
	10 25		320.00	100.00	320.00 >	449.00	590.00	180.00		320.00	207.00	73.00	9	22 00	36.00	25.60	2 20 =	22.0	0.35	0.00	0.32 -	5	2	9.	10.00	V 04	9	6.60	į	0.32	0.03	0.01	0.25	0.10	0.56	1.00 ~	13.70	2.10	9.60	0.21	6	133.00		59.80	
	10 25	6	2.50		\$2.00 >	11.20	12.10	147.00		٨	2.00		19.00	22.00	25.00	0.00	0 22	22	0.35	0.00	0.03	0.10	345	6.5	0.32	CE 0	10	5.80		0.0	0.10	0.03	0.05	0.00	0.83	1.00 <	0.00	0.10	0.21	0.32		44.30		1.50	
	Diff.	•	K :	NA.	0.995	0.590	0.626	0.918		0.03	706.0	0.70	5	•	Y V	0.758	44			6.7%	N N	**	0 041	2.0	NA NA	**	*	190		¥	¥	Y.	1.334	0.732	0.893	0.683 <		NA ~	¥	¥	603	0.559		0.575	
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	Date	767.02	00/43/01	08/01/81	68/01/50	12/05/89	12/05/89	05/30/90	00.00.00	10/11/00	10/11/02	(0/63/11	10/10/86	11/2/184	20/42/11	08/53/89	10/10/86	04/01/88	06/01/00	40/01/cn	08/03/87	08/11/87	05/10/80 214	10/01/00	10/17/86	11/25/84	05/17/88	05/10/89		10/17/86	11/25/86	08/18/87	02/07/89	08/23/89	11/01/90	04/10/90	04/22/90	10/17/86	11/05/86	08/18/87	00/36/20	12/14/89		03/06/90 NBD	
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Ship- Test Plt Diff. IC 25 11/03/86 NA 0.66 > 5 07/27/87 NA 0.66 > 5 27 09/29/87 NA 32.00 53 03/28/89 PCW 0.667 24.00 52 03/15/89 PS 0.663 4.29 52 03/15/89 PS 0.663 4.29 52 03/15/89 PS 0.663 15.30 > 62 52 03/15/99 PCW 0.868 15.30 > 62 53 08/29/89 PCW 0.868 15.30 > 62 54 03/13/90 UM 0.665 0.00 > 64 55/03/90 VM 0.775 294.00 64 05/03/90 VM 0.775 294.00 64 05/03/90 VM 0.775 294.00 64 05/03/90 VM 0.520 114.00 > 64 55/30/90 UM 0.520 114.00 > 64 55/30/90 VM 0.520 114.00 > 64 55/03/90 VM 0.809 102.00 64 05/03/90 VM 0.809 102.00 64 05/03/90 VM 0.806 4.11 64 11/01/90 ZM 0.885 < 3.20	50.57 8.34 115.00 0.00 0.00 7.69 9.72 210.00 0.00 0.00 0.00 0.00 0.00 0.00 0
Ship- Test Plt Diff. IC 25 11/03/86 NA 0.66 > 5 07/27/87 NA 0.66 > 5 27 09/29/87 NA 32.00 53 03/28/89 PCW 0.667 24.00 52 03/15/89 PS 0.663 4.29 52 03/15/89 PS 0.663 4.29 52 03/15/89 PS 0.663 15.30 > 62 52 03/15/99 PCW 0.868 15.30 > 62 53 08/29/89 PCW 0.868 15.30 > 62 54 03/13/90 UM 0.665 0.00 > 64 55/03/90 VM 0.775 294.00 64 05/03/90 VM 0.775 294.00 64 05/03/90 VM 0.775 294.00 64 05/03/90 VM 0.520 114.00 > 64 55/30/90 UM 0.520 114.00 > 64 55/30/90 VM 0.520 114.00 > 64 55/03/90 VM 0.809 102.00 64 05/03/90 VM 0.809 102.00 64 05/03/90 VM 0.806 4.11 64 11/01/90 ZM 0.885 < 3.20	77.20 77.20 83.00 100.00 74.50 74.50 21.00 21.00 21.00 21.00 21.00 21.00 21.00
Ship- Test Plt biff. 5 11/03/66 NA 5 07/27/67 NA 5 07/27/67 NA 53 03/28/69 PCU 0.667 52 03/15/69 P3S 0.663 52 03/15/69 P3S 0.663 52 03/15/99 PCU 0.667 53 08/29/89 PCU 0.667 64 03/13/90 UJX 0.656 65 04/12/90 VA4 0.775 64 05/03/90 VX5 0.809 67 05/33/90 UJY 0.520 64 05/03/90 VX5 0.809	0.28 9.50 9.50 95.70 6.60 \ 14.30 14.30 1.67 0.00 0.00 0.00 0.00 0.32 \ 3.20 10.00
Ship- Test Pit ment Date # 5 11/03/86 55 07/27/87 53 03/28/89 PCW 52 03/15/89 PCW 52 03/15/89 PCW 52 03/15/89 PCW 64 03/13/90 UJX 64 05/03/90 WK5 64 05/03/90 WK5 64 05/03/90 UJZ 64 05/03/90 VX5 64 05/03/90 VX5 64 05/03/90 VX5	0.906 0.873 0.666 0.943 1.003 1.005 0.723 0.723 NA NA NA NA NA NA NA NA NA NA NA NA NA
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		Ĭ	3.10	0.25	0.03	76 00	15.71	2 10	200	3 8	3	3.10	0.00	3.80	0.00	0.00	3.20	1.70	5.30	1.30	09.0	0.00	2.50	1.00	1.70	3.20	1.00	0.00	1.20	0.05	1.40	0.30	4.53	24.52	
	5	5	5.30	0.00	0.00	7 30	3.86	202		3 8	3	6.10	0.00	000.00	0.00	0.00	00.00	0.30	0.65	92.00	0.50	0.00	12.80	0.39	0.16	100.00	0.38	0.00	1.80	0.00	3.60	8.	0.00	5.31 > 0.00	
		8	0.00	0.00	0.00	6	88.	2	3 6	3 8	3	0.00	0.00			0.00	31.30 > 1		0.00			0.00	3.20					0.00	0.00	0.00	0.00	0.0	0.00	0.00	
		5 S	320.00	320.00	100.00	00 002	948.00	20 00	320.00	200.00	90.00	320.00	320.00	320.00	320.00	320.00	100.00	100.00	32.00	320.00	320.00	100.00	100.00	100.00	100.00	100.00	320.00	320.00	10.00	10.00	32.00	32.00	32.00	100.00 320.00	
		E E	0.00	0.00	0.00		30.0	5		-		0.00		0.00					0.00			0.00			0.00			0.00	0.00	0.00	0.00	0.00	0.00	0.00	
mt'd)	:	N 20	5.30	0.00	0.00	4	. 'Y	2 20		3 6	3	6.10	0.00	1000.00	0.00	0.00	100.00	0.30	99.0	56.00	2.00	0.00	12.80	0.39	0.16	100.00	0.38	0.00	1.80	0.00	3.60	1.00	0.00	5.31	
Table 21 (Cont'd)	1	5 5 7	100.00	100.00	100.00	200 002	485.00	220 00	220.00	00.026	37.00	320.00	100.00	320.00		320.00	32.00 >	1.00	1.00	320.00 -	320.00 ~	97.00	32.00	3.20	3.20	32.00 >	10.00	39.70	3.20	3.20	32.00	10.00	23.00	100.00 >	
10		05 21	19.00	0.00	0.00	200	6.19 6.10	02 70	200	3 6	3	52.30 >	0.00	0.32 >	0.00	0.00	0.32	3.30	1.48	5.70 -	~ 00.09	0.00	2.50	8.30	19.90	0.32	26.40	0.00	1.80	0.00	8.80	10.00	0.00	18.80 > 0.00	
		VI 23	313.00	0.31	0.00	77	7.57	21 00	3	3 6	3	20.00	0.00	> 00.0001	0.00	6.30	> 00.99	99.0	99.0	000.000	0.60	0.00	90.99	99.0	0.21	> 00.99	0.21	0.00	2.00	0.00	10.00	1.00	2.10	0.00	
		C 21	160,30	96.00	_		262.00	210 00	20.07	20.00	6.00	320.00	99.00	320.00		160.00	21.00 >	9.0	99.0	320.00 >1000.00	32.00	52.00	21.00	2.10	2.10	21.00	9.90	22.00	1.60	1.60	32.00	9.60	13.60	100.00 > 121.00	
	1	S 31	< 0.32	210.00	0.00	7 07	34.60	10 00	9	9 6	3	16.00 >	0.00	< 0.32 >	0.00	0.00	< 0.32	1.00	1.00	- 0.32 <	~ 50.00	0.00	0.32	3.20	10.00	< 0.32	32.00	0.00	< 0.32	0.00	3.20	9.90	6.50	10.00	
			¥	¥	0.739	36.	1.252	4		MA 0	3	¥	¥	¥	¥	¥	¥	4	¥	N.		0.721	¥	¥	¥	¥ ¥	¥	1.127	¥ .	¥	¥	¥	0.873	0.535	
	<u>۔</u>		:	:				;				:	:	;	:		:	:		;	:		;	;	:	;	:		;	:					
		Oate	10/22/86	05/18/88	08/30/89 RER	04/11/00	08/08/89 R29	10/24/84	06/2/100	05/54/88	21 15/ 22	11/12/86	05/24/88	10/31/86	05/04/88	06/01/88	10/31/86	07/27/87	05/24/88	10/31/86	05/24/88	04/11/89	11/12/86	07/28/87	05/25/88	11/12/86	07/27/87	05/23/89 064	11/12/66	12/02/80	11/13/86	05/25/88	08/30/89 REU	08/30/89 REV 05/31/90 WHX	
	Ship-		~	2	2		28		۱ ۸	1 M	1	m	M	6	6	0	•	6	٥	٥	6	53	5	9	5	5	10	10	5	9	0	10	10	12 67	
		ė		1987	1987		1992	1001				2013		2159		2159	2160				2912				2219	2220		2220	2221	2221			2226	2274	
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	1796	Ħ	E !	Ē	CPE	CPE	CPE	9	9	HT	Ë	Ē	8	30	TTH	H	E	ĒĒ	ě	CPE	118	T T	Ē	TTM	ĒĒ		H H	Ē	ļ			F	TTM	TTM	E E	Ē
	Σ	72.62	35.74	9	1.10	0.00	4.60	2.50	2.60	0.15	5.6 8.6	13.73	3.50	3.40	54.10	19.21	10.59	7.40	2.40	0.0	27 5.8	11.18	0.0	31.95	0.00		8.7 5.7	8.82	3		000	11.48	9.61	17.02	7.43	3
	15	6.73	10.53 >	<u>.</u>	17.10	0.00	10.00	3.10	3.90	0.00	2.03	3.45	69.00	4.30	< 00.65	3.63 >	8.0	88.	100.00	0.00	× 07 S	0.00	0.00	9.55 >	0.00	!	1.03	3.5		27.2	0.00	2.77	2.28 >	4.00.4	8.8	3
	N 95	3.69	0.0	3	0.0	0.00	320.00	10.00	10.00	0.00	8.6	80.	0.00	100.00	23.00	0.0	8.6	8 8			9	800	0.00	0.00	0.00		0.0	9.0		38	0.00	0.00	0.00	0.00	9.6	5.0
	25 28	320.00 >	320.00	350.00	320.00	320.00	320.00 >	10.00	320.00 <	320.00	8.8	0.0	320.00	320.00	320.00 >	320.00	274.00	320.00 320.00	320.00	320.00	100	278.00	251.00	320.00	320.00		2.5	62.20	200	10.00	10.00	10.00	32.00	10.00	320.00	350.00
	10 %	86.70 >	8.0	3.5	0.00	0.00	1.00 \$	1.00 \$	32.00 >	0.00	8.6	0.00	0.00	3.20	13.90 >	0.00	8.6	88			5	800	0.00	0.00	0.0 0.0		9.5	0.0		88	0.00	0.00	0.00	0.00	9.0	3.5
nt'd)	A1 50	12.70	15.20	۲.43	17.10	0.0	1000.00	31.00	39.00	0.00	20.03	15.27	152.00	400.00	97.80	8.07	8.8	88	100.00	0.00	2	00.0	0.00	30.55	0.00		5.19	2.73	8	1.00	0.00	10.45	4.71	2.4	0.0	2.0
Table 21 (Cont'd)	10 50	326.00	259.00	M. C27	320.00	100.00	320.00	10.00 >	320.00 >	2.90	5.6 8.5	8.64	320.00	5				2 S. S.		320.00	01 29	200	17.90	320.00 >	320.00		16.10	19.60		7.5	3.07	10.00	4.22	10.00 >	100.00	3.₹
Tel	10 50	25.80	17.10	3.5	18.70	0.00	0.32 >	0.32 >	8.30 >	0.00	0.20	0.57	2.10 ~	0.80	39.1	19.30	0.0	8 8	9	0.00	2	0.00	0.00	10.50 >	0.0 0.0		3.11	7.20		3.6	0.0	0.96	0.89	2.16 >	8.6	3.0
	AI 25	11.60	25.52	2.0	9.90	0.00	10.00 <	2.00	9.00	9.	07.0	8.9	313.00	10.00	81.50	20.08	69.6	90.63	206.00	0.0	21 21	8	0.36	23.50	7.31		5.09	2.73		9.4	0.03	4.79	3.79	6.88	3.45	0.84
	10 25	174.00	180.00	149.00	210.00	66.00	3.20 >	1.00 -	32.00 ~	1.00 -	2.03	: : :	100.00	3.20	\$1.50 >	70.10 >	8:5	100.00		210.00	01 67	55.00	6.98	100.00	20.70		3.20	12.10		8 8	0.05	59.2	2.02	8.83	26.20	Ž.
	10 25	15.10	7.6	24.20	32.00	0.00	0.32				0.54	0.28	0.32	0.32			3.20	38.	0.32	0.00	1 20	27.70	19.30	4.25	26.10		1.53	4.44		× 27 C	0.83	0.55	0.53	1.28	7.59	17.0
	Diff.		0.535	0.836	¥	ž	Y Y		¥		0.652	0.908	V V	¥	0.598 <		0.80	0.493	ž	¥.	0.44.0	767.0	0.848	0.660	0.880		1.152	0.885		121	000	0.811	0.770	1.058	0.534	0.800
-	*			5	:	:	:	:	:		Z .			-	PG1			× ×					XX		VA7		192	× ×	3	3				RGK	¥2	٨٧٥
1	Date	04/11/89	08/30/89	08/18/50	12/15/86	08/18/87	12/15/86	07/28/87	05/25/88	04/11/89	08/30/89	07/12/90	12/15/86	07/28/87	04/11/89	08/30/89	04/12/90	07/05/90	12/15/86	08/18/87	08/47/80	03/13/90	05/03/90	08/30/89	04/12/90		09/06/89	05/03/90		04/11/69	05/31/90	07/12/90	10/04/90	68/90/60	03/13/90	U2/CU/CU
	ment .	53	2;	\	13	Į.	13	13	13	53	5 t	67	13	13				6 S	*	13	ž.	2 3	3	17	65	}		8 8		3 2	25	29	29		3 :	
97.6		22.75			2317	2317	2318					2318	2320				2320		2325			2363			2633		2453			Sign		2503			2206	

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1	Type	8		E	Ę	H	H	MTT	Ē	TIM	Ħ	TTM	MTT	MTT	HTT	HTT	TH	MTT	MTT	GPE	CPE	MTT	TTM	TH	MTT	CPE	MTT	E !		G	MTT	H	HTT	HTT	MTT	i i
	TAI	2.60	15 10	0	00.0	3.00	0.00	10 81	0.0	2.67	3.03	0.00	3.70	21.49	11.37	0.00	12.67	3.85	0.00	2.10	09.0	6.59	15.12	7.0	2.91	. N	14.98	32.54	6.51	3.50	0.0	12.88	11.92	5.73	7.24	0.00
	IS	13.00	V 65 0	800	0.00	0.00	0.00	4 01 0	0.00	0.02 >	0.00	0.00	0.06 ×	4 49.9	1.82 >	0.00	1.11	0.00	0.00	4.00	1.40	0.00	7.6	0.85 >	0.00	2.70	0.00	8.83 v	3.5	9.1	0.0	80.4	1.02 >	0.00	1.11	0.00
	AI 95	3.20		88	0.00	0.00	0.00	8	0.00	0.00	0.00	0.0	0.00	0.00	0.00	0.00	0.00	0.00	0.00	3.20	0.00	0.00	0.00	0.00	0.00	10.00	0.00	8.6	3.	1000.00	0.00	0.0	0.00	0.0	0.00	9.0
	70 %	32.00	2 20	35.00	6.11	3.20	9.43	320.00	320.00	320.00	1000.00	320.00	1000.00	320.00	320.00	1000.00	0.01	0.10	0.32	320.00	100.00	320.00	30.90	66.30	25.60	32.00	3.20	9.0	3.20	320.00 >	320.00	3.20	78.80	88.20	320.00	320.00
	10.98	10.00	200	× 000	0.00	0.00	0.00	* 00 0	0.00	0.00	0.00		0.00	0.00	0.00	0.00	0.00	0.00	0.00	100.001	0.00	0.00	0.00	0.00	0.00	3.20	0.00	0.0	8.5	0.32	0.00	0.00	0.00	0.00	0.00	0.00
mt/d)	AI 50	70.00	77 0	8	00.0	0.00	0.00	9	0.00	1.68	0.00	0.00	2.31	12.51	1.82	0.00	1.11	0.0	0.00	13.00	1.40	0.00	2.50	1.54	0.0	27.00	0.0	15.60	3	000.00	0.0	87.05	1.9	0.00	1.11	0.0
Table 21 (Cont'd)	1C 50	32.00	3 20 >	8	50.	3.20	96.0	320.00 >	320.00	320.00 >	858.00	320.00	1000.00	320.00 >	320.00 >	1000.00	0.01	0.03	0.32	320.00	100.001	93.00	4.48	5.85	3.10	32.00	3.20	3.09	70.7	320.00 >	1.0	3.20 >	19.50	21.40	320.00 >	JOS.00
T.	10 50	08.0	S 02 >	0.0	0.00	00.0	0.00	× 05 09	0.00	191.00 >	0.00	٨	434.00 >			0.00	• 10.0	0.00	0.00	25.00 >	4 00.02	0.00	1.73	3.77	0.00	1.20	0.00	0.5 0.6	3.	0.32	0.00	0.06	10.00	0.00	289.00 >	0.00
	AI 25	31.00	7 07	0	0.00	0.00	0.00	41 0	0.00	0.05	2.15	0.00	0.13	11.62	2.96	0.00	2.56	0.95	0.00	2.00	5.00	0.00	18.60	3.20	1.61	9.00	5.20	54.73	77.7	1.00 <	0.0	8.9	5.49	1.32	2.12	9.0
	10 25	10.00	2 62	2 2	0.58	3.20	0.52	80 4	320.00	3.06	462.00	320.00	25.70			3.20	0.01	0.05	0.32	100.001	100.001	25	1.86	3.20 >	1.82	3.20 ~	2.19	۲. ۲.	6	0.32 >	1.0	97.0	10.20	11.20	320.00 >	
	10 25	0.32	2	00.00	0.00	00.00	0.00	17 00	0.00	62.20	215.00	0.00	195.00	14.60	108.00 >	0.00	0.00	0.05	0.00	~ 20.00	~ 50.00	0.00	0.10	1.00	1.13	~ 0.50	0.42	0.03	8	< 0.32	0.00	0.0	4.11	8.50	151.00 >	0.00
	Diff.	¥	1 288	1.118	1.090	709.0	1.030	1,103	0.740	1,190	0.834	1.190	0.927	1.159	1.169	956	1.352	1.016	0.808	× ×		0.601			0.643	Y.	1.141	0.725	0.710		0.727	0.677	0.700	0.702	1.064	0.747
2	*	:	3			SS	ZYA	059	108	RGP	VAB	RGP	VAC	RGO	VEH	X87	000	SPV	9	:	:	P62	OK3	PNB	E E	:	*	E I	ž	:	PG3	Z.	PAT	PNC	VEY	68 80
	Date	02/09/88	02/07/80	05/23/89	09/06/89	12/05/89	11/01/90	09/07/00	12/05/90	68/90/60	04/12/90	68/90/60	04/12/90	68/90/60	04/11/90	02/03/90	02/07/89	11/29/89	12/02/90	09/14/87	05/25/88	04/11/89	02/07/89	04/11/89	09/01/89	09/01/87	02/07/89	09/02/89	40/63/11	02/02/88	04/11/89	09/01/89	04/04/89	04/19/89	04/11/90	07/03/70
	Salp-	22	87	5	21	87	15/51	10	10	4		19			65		87	61	19	92	56	53	87	27	52	22	87	55:	5	28	53	28	32		\$9	
	NO.	2563						2586	2586	2590			1652			2631		2812				5906				2980		2980				3038	3450		3491	

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Assay	Type	Ħ				ĒĒ	ĒĒ		EE		ĒĒ	ĒĒ	ĒĒ	FFFFF
	¥	0.00	25.91 15.48 1.71	41.57 7.98 13.61 31.92 2.14		33.34	31.82	2.93 37.07 0.41 44.49	3.04	19.88 10.22 0.00 8.11	2.22	1.81	9.14	4.54 0.00 10.28 5.36
	2	3.39	4.32 > 0.00 >	0.00	3.20	5.91 0.00	8.33 >	1.75 × 5.51 × 0.03 × 54.37 ×	0.58 >	4.81 2.09 0.00 1.07 v	0.00	0.00 >	1.43 \	0.00 0.00 0.00 0.00 0.00 0.00 0.00 0.0
	VI 95	0.00	0.00	000000	0.00	0.0	3.36	33.49	0.00	8888	0.00	0.00	0.00	00000
	26 27	326.00	100.00 320.00 320.00	320.00 320.00 320.00 1000.00 320.00	320.00	320.00 942.00	320.00 × 939.00	228.00 10.00 v 10.00	32.00	318.00 306.00 259.00 304.00	320.00 944.00	320.00	320.00	9.20 9.72 9.94 9.94
	10 95	9.0	0.00	00000	0.00	88.	0.00	0.00	0.00	0.0.00	0.00	0.00	0.00	00000
wt,d)	A1 50	6.07	5.59 6.03 0.00	12.20 0.00 0.00 0.00		0.00	5.71	3.20 62.45 0.14 54.37	19.20	3.80 3.55 0.00 2.05	0.00	9.00	1.43	0.00
Table 21 (Cont'd)	10 50	182.00	100.00 × 320.00 × 320.00	320.00 × 320.00 × 1000.00 × 320.00		320.00 >	320.00 >	3.20 v 10.00 v 0.69 10.00 v	19.20 >	175.00 175.00 163.00	320.00	320.00	320.00 > 782.00	2.16 5.35 6.56 6.42
Į.	10 50	30.00	17.90 > 53.10 > 0.00 >	26.10 × 0.00 × 0.00 × 205.00 × 0.00 ×	0.00	42.50 v	32.80 >	1.00 0.16 > 4.87 0.18 >	1.00	19.90 49.30 0.00 79.60	291.00	0.00 >	223.00 > 0.00	88888
	A1 25	5.88 0.00	7.14 4.85 1.73	42.50 0.83 3.52 117.68 1.98	0.00	0.00	20.40	1.75 7.79 0.84 76.87	0.58 < 1.21	7.48 5.05 0.00 1.80	1.48	0.00	2.43	1.58 0.00 3.47 2.56
	TC 25	102.00	77.30 75.70 32.00	320.00 × 266.00 × 320.00 × 1000.0 × 252.00	181.00	251.00	273.00	0.88	0.58 >	95.70 103.00 51.60 85.20	271.00	320.00	320.00 >	1.58 3.78 4.81 6.63
	IC 25	0.00	10.80 15.60 18.50	7.52 × 320.00 90.90 × 8.50 × 128.00	1.94	0.00	13.40	0.11 0.18 0.13 v	^ 1.86	12.80 20.40 0.00 47.40	183.00	0.00 >	132.00 > 207.00	0.00
	Diff.	0.732	1.270 0.764 0.803	0.509 0.680 0.741 1.016 0.792	0.529	0.531	1.011	0.623 0.786 0.903 0.811	1.041	0.519 0.704 0.866 0.747	0.866	0.742	0.742	0.679 0.903 0.747 0.779 0.720
	Date	03/29/89 PF6 09/07/89 RIL	11/03/88 02A 04/19/89 PNF 09/12/89 RK3	01/11/89 OAE 03/07/89 OXA 09/12/89 RK4 11/29/89 SPV 04/24/90 VL7	11/08/88 0C5 12/20/89 100	11/09/88 0c9 12/20/89 10P	11/10/88 OCP 12/14/89 SXW	12/13/88 061 09/12/89 RKS 05/31/90 WI1 07/12/90 XHD	12/13/88 061 10/11/89 RT2	12/13/88 062 09/12/89 RK7 04/24/90 VLA 07/03/90 XB9	04/24/90 VLA 07/03/90 XBA	04/24/90 VLB 07/06/90 XDB	04/24/90 VLB 07/06/90 XD9	09/12/89 RK9 05/31/90 W11 10/02/90 28L 10/04/90 ZAC 10/04/90 ZAC
	No. ment	88 82 82	222	92 ABEBE 92 51 92 51 92 61		32	22 22 28 32 32 32	67 38 87 38 87 38 87 38	33 35 33 35	****	\$ \$ 8 \$ 22	55 65 55 65	8 8 8 8	12 27 27 27 27 27 27 27 27 27 27 27 27 27
3	*	3558 3558	3584 3584 3584	3592 3592 3592 3592 3592	3615 3615	3621 3621	3688	3802 3802 3802 3802	3803	3935 3935 3935 3935	4032	4035	4036 4036	4051 4051 4051 4051 4051

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	Assay			H	CPE	CPE	FE	CPE		H	FE	H	EE	E E	FF	FE
	ĭ	33.61 1.50 22.05	6.49	8.28	1.30	1.10	43.67	9.97	27.00 20.88 0.00	36.13 15.52	27.07	39.64	50.81	43.60	40.66	10.37
	ï	10.47 > 0.00	0.00	3.98 ×	1.60	1.20	9.24 >	1.50	4.92 v 2.06 v 0.00	4.13 > 0.00 >	1.76 >	12.20	3.20 >	20.00 >	19.00 >	80.0
	A1 95	3.31	26.50 >	10.30	3.10	3.20	26.00	0.00	0.00	3.73	0.00	11.80	0.00	0.00	0.00	0.00
	70 95	1000.00 > 959.00 1000.00	0.03 26.50 > 10.00	92.30 30.90	100.00	320.00	32.00	320.00	100.00 100.00 320.00	30.70	100.00	320.00 >	320.00 >	320.00	320.00	261.00
	10 95	302.00 >	0.00	9.60	32.00	100.00	2.91	0.00	0.00	8.22.	0.00	27.20 > 21.10 >	192.00 >	0.00	0.00	0.00
nt'd)	A1 50	13.44 0.00 5.55	0.00 2.77 < 0.00	8.18 7.70	9.30	0.00	15.90	54.00	2.92	0.00	6.29	31.30	19.80	5.50	3.20	3.20
Table 21 (Cont'd)	10 50	1000.00 × 585.00 679.00	0.03 2.77 × 1.94	28.50	100.00 ~	320.00 ~ 100.00	19.90	320.00 - 63.00	100.00 > 100.00 > 267.00	6.60	48.40	219.00	320.00 >	320.00 >	320.00	54.00
Tak	10 50	74.40 > 10.00	0.00	3.48	6.30 ~ 2.50	26.00 ~ 3	1.25	5.90 -	20.30 > 48.40 > 0.00	1.87 00.00	11.30	7.02	16.20 >	16.00 >	17.00 >	17.00
	AI 25	34.29 1.58 8.70	0.00	7.86	3.10	1.60	18.40	3.00	8.05 7.38 0.00	9.76	4.05	23.10 193.22	78.30 31.86	40.00 56.00	31.00	3.40
	10 25	779.00 378.00 492.00	0.03 1.89 > 0.85	13.90	10.00	32.00 -	11.60	10.00 ~	100.00 > 100.00 >	7.58	19.80	193.00 >	320.00 >	320.00 >	320.00	17.00
	10.25	22.73 239.00 56.60	1.00 • 0.69	1.76	3.20	20.00	0.63	3.20	12.40 × 13.50 × 0.00	0.78	3.80	3.73	4.00.01	8.00 >	10.00	5.00
	Oiff.	0.622 0.862 0.646	1.222 0.524 < 0.655	0.727	NA -	NA - 0.721	0.810	1.002	0.580 0.734 0.531	0.615	0.615	1.225	0.866	0.689	0.689	0.751
	= *	SSE TFF TM3	SUR UOD	PG3 RKC	 0F2	: #	PFE RT3	96	RT3	PCV RT4	PCW RT4	R 70	022 RT5	03S RT5	03S R16	031 RT6
	Test Oate	12/05/89 01/18/90 02/08/90	02/07/89 12/07/89 02/13/90	04/11/89	05/11/88 01/24/89	10/05/88 03/29/89	03/29/89	10/12/88 03/29/89 PFF	04/05/89 10/11/89 12/20/89	04/05/89	04/05/89	07/06/89 08/16/89	12/06/88	12/07/88	12/07/88	12/07/88
	Ship- ment	333	3 3 3	22.23	6 3	33	23	33	333	33	33	22	45	55	45	45
	AVS	1207	7207	7540	4241	4420	4427	4428	4432 4432 4432	8577	6577	4452	5977	477	£73 £73	22 23

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Assay	178	ĒĒ		E	H		HTT	HT	ĒĒ	***		Ē	I	Ħ	HT	HT	FI	H			E	E	Ē	EE	EE	Ę		ĒĒ	EE	
	IAI	30.17	87.8	0.0	0.03	2.33	7.46	13.40	2.22 0.10		5			54.23		20.12	16.01	10.58	4.21	5	2.5	30.00	?	 22	35.19	16.63		41.63	5.20	
	15	1.00	0.87 >	0.0	0.0	0.00	1.45	1.41 >	9.0		38	0.0	4.85 >	62.74 >	0.00	4.04	2.88	0.0	8 8		83	5.2	3	0.00	6.89	3.43 >	6.63	3.49	0.0	
	N %	0.0	00.0	8.0	8.0	0.00	0.00	8.8	88	8	3 2	0.0	0.00	3.61	0.00	0.0	0.00	8.6	8 8	8	88	3.52	3	88	30.30 >	3.27	3	0.00	9.0	
	٦ گ	320.00	97.30	100.00	9.00	100.00	100.00	100.00	82.29 82.30	8 87	5 6	0.92	320.00	1000.00	1000.00	3200.00	96.90	2.50	87.18 5.10	8	8.1 0	3.8	2.40	32.00 92.90	30,30 >	9.6	64.66	320.00 >	320.00	
	5 8	0.00	00.0	0.00	0.00	0.00	0.00	8. 8.	88.	5	86	0.00		277.00 >	. A	0.00	0.00	0.0	8 8	8	88	27.50	3	88	1.00	2.95	3	222.00 > 0.00	0.00	
(p, y	VI 20	0.00	2.12	0.0	8.8	8.0	2.83	3.15	8.8	1 87	5 5	0.0	4.85	. 69.79	0.00	2.66	4.15	8.6	8 8	2	0.0	9.68	3	- ° 28	15.10 <	3:	3.36	6.39	7.37	
Table 21 (Cont'd)	05 22	320.00 >	17.60	100.00	8.8	35.00	78.70	67.50	67.6	9	3,5	0.25	320.00 >	1000.00	000.000	3200.00 >	62.60	45.20	51.40	97 27	24.80	8.8	3.0	31.00	15.10 >	6.60	8.5	320.00 >	25.70	
•	10 50	18.00 >	8.27	0.00	8.0	0.00	27.80	21.50	88	5		0.0	< 00.99	15.50 > 1	0.00 > 1	566.00 > 3	15.10	8.6	8 8	0 0	9.0	3. ce	3	5.31 0.00	38.0	1.43	8.5	50.10 > 140.00	28.50	
	AI 25	19.00	3,56	0.0	0.0	0.00	2.63	3.32	0.00	8	88	2.24	7.62	143.30		8.38	2.67	9.15	3 %	8	0.0	16.00	9.0	0.00	8.06 < 12.96	8:		16.50 13.93	9.00	
	22 21	198.00	7.20	100.00	90.00	21.50	40.30	30.40	85.9 8.38	20.0	0.18	0.14	320.00 >	970.00	477.00	2290.0	43.50	18.40	25.50	27	17.20	66.64	2	5.94 18.20	8.06 >	6.9	67.0	320.00 >	2.47	
	S 21	10.00		0.00	A		15.30	9.16	0.00	200		90.0	45.00 >		208.00				13.08 8.08		0.00	. s.	2.	0.00	0.73	9.9	8	19.40 > 35.20	2.94	
	oiff.	25.0	1.259	0.729	0.690	0.534	0.749	0.859	0.655	0 785 c		0.713		0.857	0.595	0.533			0.850		0.658	0.853		0.971	0.857 <			1.098	1.114	
Test Pit	Date	12/07/86 03V 10/11/89 RT7	02/01/89 out		02/13/90 UOR	1)3/30 WOL		11/29/89 SPX	02/13/90 UOM	CV0 08/10/20	12/05/89 SSI		4/24/90 VLJ	80X 06/90/20	12/05/89 SSF	02/08/90 TM	000 68/90/20		05/08/90 UFO 05/03/90 VX9		12/05/89 SSK	11/01/88 00L	Z/ 14/07 SAA	11/01/88 00N 03/06/90 UDV	11/01/88 000 12/14/89 sxv	11/02/88 010	12/20/09 100	11/02/88 01F 12/14/89 SXT	11/02/88 01H 12/07/89 sus	
Ship	ment.	\$\$ \$\$		0 27			97		3		87		65 04			GABSH O			88		**	7,		22	33	3:		33	22	
AVS	<u>.</u>	4524	4527	4527	4527	1754	4592	4592	4592	7400	0097	6094	4611	4611	4617	4617	123	627	£136	7727	77.27	4747		125	4753	424	*()*	137	4762	

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100	1ype	THE	EE	EE	ĒĒ	EE	EE	EE	EE	EE	EE	EEE	EEE	EE	EE	EE	EE	
	¥	52.59 13.63	27.25	12.82	19.77	1.88	24.98	4.38	62.20 53.61	42.28 51.00	19.23	37.03 59.59 42.18	23.35 23.39 24.34	33.29	2.72	0.00	18.70	
	15	0.37 >	3.02 >	0.00	3.43 >	3.79 >	4.00.0 0.00	7.20	%.S.	23.80 >	1.33 \$	6.23 21.49 > 7.98 >	10.90 6.82 45.46 >	4.93 v	1.8	0.00	3.01	
	AI 95	0.00	0.00	0.00	0.0	0.00	0.00	0.00	39.50	39.84	0.00	0.00	34.04	0.00	0.00	0.00	0.00	
	25 25	320.00 >	320.00	306.00	302.00	296.00	306.00	320.00	320.00 >	320.00	320.00	320.00 320.00 >	320.00 320.00 1000.00 >	320.00	8.52	320.00	333.00	
	26 52	0.00 0	0.00	0.0	0.00	0.00	0.00	0.00	8.10 > 25.40 >	8.03 ×	0.00 \$	29.60 \$ 0.00 \$	0.00 \$	0.00	0.00	0.00	0.00	
(p,1	AI 50	2.39	3.02	2.03	8.15 0.00	6.41	0.00	25.80	44.00	50.00	1.33	11.30 21.49 7.98	15.20 14.50 73.50	6.93	0.00	 80.	5.51	
Table 21 (Cont'd)	10 50	320.00 >	320.00 > 206.00	181.00 28.10	137.00	92.40 183.00	179.00	25.80 >	320.00 > 1	222.00 76.20	320.00 > 767.00	221.00 320.00 >	320.00 > 1000.00 >	320.00 >	2.21 > 0.66	284.00	164.00	
Tab	10 50	6.40 > 3	106.00 \$ 0.00	99.20	0.00	14.40	0.00	1.00	2.22 > 5.59	2.11	241.00 >	19.50 14.90 > 12.50 >	21.10 > 22.10 > 13.60 > 1	0.00 • 1	0.00	0.00	33.30	
	AI 25	1.06	0.00	0.00	11.30	18.00	25.90	7.20 < 2.06	141.00	83.70	2.54	16.80 31.51 84.69	15.90 11.10 98.58	320.00	1.29	2.10	5.55	
	25 21	26.70	320.00 > 121.00	9.24	57.50 140.00	54.70	108.00	7.20 >	210.00	106.00 54.10	320.00 > 544.00	122.00 320.00 >	230.00 151.00 619.00	320.00 > 159.00	1.29 > 0.49	147.00	100.00	
	10 25	46.80	4.43 >	2.01	5.10	3.04	4.18	1.00	3.04	1.26	126.00 >	7.22 10.20 > 1.18 >	14.50 13.60 6.26	1.00 \$	0.00	0.00	18.10	
	Diff.	1.054	0.827	0.845	0.967	0.914	0.914	0.684 <	1.021	1.057	1.221	0.808 0.831 1.070	0.418 1.167 6.873	1.142 < 0.489	1.265 < 0.489	1.131	1.166	
2	*	100	100	0C1	0C2	003 004	500	270	SPY	00G SY1	SP2	201 105	S 20 28	100	108	OHO	005	
	Date	11/02/8 6 011 01/10/90 100	11/08/88 12/20/89	11/08/88 02/13/90	11/08/88 02/13/90	11/08/88 02/13/90	11/08/88 03/06/90	01/12/89	01/12/89	02/01/89	02/14/89	02/22/89 11/29/89 12/05/90	01/17/89 01/24/89 11/29/89	01/18/89	01/18/89	01/31/89	02/01/89 11/29/89	
	ment .	33	33	33	33	33	33	99	97	87	848	84 64 84	995	99	99	979	97	
	0	4763	178	4765	4768	6927	£139 £139	4785	9627	4822	4827	4855 4855 4855	4871 4871 4871	4875 4875	1687	1167	4919	

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	Assay Type	EE	FF	ĒĒ	FE	EEE	EEE	EEE	EEE		EE	EEEE	EEE
	TAI	36.63	37.32 24.80	16.9 0.60	10.53	31.50 1.46 3.74	3.3 3.3 3.4	26.19 2.52 0.00	46.17 12.52 0.60	19.88 4.91 7.6.7	0.00	22.70 23.52 19.08 52.11	6.42 13.45 24.40
	15	15.40 >	10.80 >	173.00 \$	151.00 > 0.00 >	14.50 v 0.00 0.00	16.00 v 0.00 v 141.00 v	4.52 0.00 0.00	6.36 0.00 0.00 0.00	36.60 0.00 0.00 0.00 0.00 0.00	320.00 >	5.66 v 6.85 v 28.63 v	0.00 v 2.14 v 4.79 v
	A1 95	0.00	0.00	0.0	0.00	000	0000	198.00 0.00 0.00	0.00	00.00	A .	0.00	0.00
	8 21	30.40	30.30	320.00 320.00	320.00 320.00	320.00 320.00 100.00	328.00 309.00 320.00	198.00 × 92.40 10.00	340.00 311.00 320.00	320.00 320.00 305.00	320.00	320.00 1000.00 1000.00 3200.00 >	320.00 1000.00 1000.00
	S 21	2.98	0.00	9.0	0.00	888	888	0.00	31.60 0.00 0.00 v	98 88		0.00 v 1 0.0	0.00
(P)+		22.60	20.60	173.00	278.00	28.70 0.00 0.00	184.00 0.00 141.00	8.03 0.00 0.00	11.30 0.00 0.00	65.30 0.00 6.65	320.00	5.66 12.26 9.96 41.22	0.00 2.14 4.79
Table 21 (Cont'd)	TC 50	22.60 16.30	20.60 > 21.00	320.00 > 1	278.00 > 2 241.00	228.00 95.00 77.90	184.00 × 1 210.00 320.00 × 1	8.03 × 10.00	191.00 211.00 320.00	202.00 225.00 52.90	A	320.00 > 1000.00 > 737.00 > 3200.00 >	320.00 1000.00 >
-	05 21	1.00	1.00	1.85 v	8.0	0.00	2.28	90.0	16.90 0.00 0.00 v	3.09 0.00 0.00 0.00	A A	56.66 × 12.66 × 17.60 × 3	0.00 × 468.00 × 10 × 10 × 10 × 10 × 10 × 10 × 10 ×
	AI 25	15.40	10.80 <	312.00	151:00 < 0.00	26.00 0.08 0.00	104.00 × 0.00 286.00	4.52 < 0.00 0.00	73.00 2.35 0.00	86.10 0.00 19.50	320.00 < 0.00	12.40 21.20 9.51 %.70	2.02 7.33 8.67
	70 25	15.40 > 8.69	10.80 × 15.50	320.00 > 3	151.00 > 1 150.00	115.00 63.50 55.00	104.00 > 1 155.00 320.00 > 2	4.52 × 5.35 0.09	108.00 156.00 320.00	113.00 163.00 19.50 >	A	320.00 > 559.00 463.00 2220.0	320.00 > 1000.0 > 1000.0 >
	82 33	1.80	8.8	1.02 > 3	9.0	2.50 0.00 1.00 1.00	1.08	80.0	1.47 66.10 0.00 > 3	1.31	0.00	25.70 × 3.50 × 3.	158.00 > 3 136.00 > 1 115.00 > 1
	Diff.	0.836 < 0.506	0.842 < 0.506	0.187	0.203 <	0.226 1.194 0.480	0.182 < 1.359 0.182	0.155 < 1.282 0.594	0.155 1.282 1.053	0.215 1.229 0.232 <	0.182 < 1.289	0.853 0.595 0.572 0.533	0.853
	==	SXC	SXC	25	P26	75 P25	2 5 E	PZ SXT SXT	25 A 9	28 82 28 25		040 SSF 10S	386 75
	Test	03/07/89 OXC 12/14/89 SXU	03/07/89	03/14/89	03/14/89	03/14/89 04/20/89 12/14/89	03/14/89 04/20/89 03/14/89	03/14/89 04/20/89 12/14/89	03/14/89 P2I 04/20/89 PUR 01/10/90 T0P	03/14/89 04/20/89 03/14/89 04/20/89	03/14/89	12/07/86 12/05/89 12/20/89 02/08/90	12/07/88 12/05/89 02/08/90
	Ship-				2.2	25.5	255	222	222	22 22	2.5	45 62 45 648N	45 62 648 84
	5	2.2	22	22	22	SON	SON	N IN IN	SON	NN NN	N N	4 4 4 4	232

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	Assay	<u>×</u>	TTH	TH	HTT	TH	TTH	HT	HT	H	TH	HTT	HTT	EE	***		E	H	TH	HT	TTH	HTT	TTM	TH	TTH	H	HTT	H	HTT		Ē	FF	1	E E	FF	
	;	ž	31.02	44.20	12.93	33.40	45.00	06.67	4.18	21.74	0.00	0.00	25.11	1.65 22.15	¥ 9	35	35.00	2.03	0.52	4.57	0.01	4.83	1.23	6.30	0.00	7.0%	20.48	48.27	13.39	40.15	3	0.24 16.64	76 2	1.46	0.00	
	i	7	6.37 >	8.67 >	2.8	6.79	20.20 >	24.62 >	0.00	62.40 >	0.00	0.00	7.21 >	7.50 \$	07.01	00.00	9.43 >	0.00	0.00	1.02	0.00	1.50	0.00	2.0	0.00	1.69	4.95	13.44 >	3.16 >	A 95.0	3	0.00	0 0	0.00	0.00	
	8	2	0.00	3.46 >	0.0	3.38 ×	0.00	27.9	0.00	0.00	0.00	0.00	0.0	0.0		0.0	0.0	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.0	0.00	6.97	0.00	3.5	3	0.00	8	88	0.00	
		ت ر	320.00	4 00.000	00.00	000.00	320.00	< 00.000	000.00	284.00	294.00	268.00	320.00	1000.00 320.00	200 002	000000	320.00	1000.00	320.00	944.00	320.00	30.60	21.00	8.8	94.80	8.3	308.00	309.00	320.00	80.00	00.00	287.00	205	294.00	9.29	
	2	2	0.00	A	0.00	A		^	A	0.00	0.00	0.00		0.00				0.00	0.00	0.0	0.00	0.00	0.00	0.00	0.00	0.00	0.00	44.30	0.00	,	A .	0.00	2	88	0.00	
1t'd)	9	N 30	6.37	8.67	3.47	2.9	20.20	26.56	0.00	142.00	0.00	0.0	14.30	0.00 10.27	07 04	00.0	10.00	0.00	0.00	.	0.00	2.41	00.00	1.17	0.00	2.57	7.01	18.21	3.44	25.00	8.5	3.43	*	0.00	0.00	
Table 21 (Cont'd)		2	320.00 >	• 00.000	^ 00.000	000.000	320.00 >	< 00.000	1000.00	142.00 >	91.60	57.10	279.00	759.00 320.00 ×	200 002	502.00 502.00	320.00 >	745.00	320.00	443.00	11.30	18.30	86.9	8.91	35.90	29.90	202.00	210.00	320.00 >	678.00	20.00	78.60	00 000	89.70	2.98	
1		1C 20	50.30 >	A	A	A		٨	٨	9.	0.00	0.00	19.50	31.10 ×	46.60	00.0	32.00 >	0.00	0.00	227.00	0.00	7.57	0.00	7.62	0.00	23.30	28.80	11.50	93.00 >	25.70	3	0.00	04 20	80.0	0.00	
		0 ¥	12.10	21.37	4.92	10.00	33.80	76.99	3.23	62.40 <	0.00	0.00	11.60	1.03	9	0.40	25.70	0.00	1.26	1.81	95.0	3.01	0.00	4.12	0.45	3.27	8.40	36.85	5.80	13.03	*	3.50	4 70	0.00	5.01	
	30	9	320.00 >	1000.0 >	835.00	1000.0 >	320.00 >	927.00	527.00	62.40 >	61.80	25.00	141.00	265.00 234.00	230 00	100.00	302.00	391.00	204.00	232.00	99.4	11.40	8.09	5.51	11.80	39.40	143.00	155.00	294.00	00.64	00.010	47.10	8	52.70	1.00 -	
		2	26.40 >	* 08.9	170.00	100.00	4 27.6	13.80	163.00		0.00	0.00	12.10	15.80	8	200	11.70	0.00	162.00	129.00	8.27	3.78	0.00	1.34	26.60	12.00	17.00	4.21	50.70	38.30	DO 7.700	61.40	20	0.00	. 1.00 < 0.25	
	777		0.907	0.570	0.572	0.589	9.864	0.559	0.570	0.982 <	0.907	0.818	0.743	0.907	27.7	0.905	0.895	0.826	1.278	1.430	1.104	0.809	1.072	0.841	1.394	0.868	1.318	0.536	1.318	0.536	00.0	1.331	171 1	0.834	1.058 < 0.581	
	Pit		110	286	108	3	3	SY2	101	9	RVC	VN V	200	RVC	5	2	RSX	VNS	990			RSV	N CO	RSX		R62	8			2 3		870 870	5	55	R65	
	Test		02/28/89	12/05/89	12/20/89	02/08/90	03/01/89	12/14/89	12/20/89	03/01/89	10/03/89	04/26/90	03/01/89	10/03/89	04/10/20	10/03/89	10/11/89	04/56/90	06/20/89	08/08/89	06/21/89	08/15/89	06/27/89	08/15/89	07/11/89	08/15/89	07/11/89	08/16/89	07/11/89	08/16/89	DK/02/50	07/11/89 08/16/89	07/11/80	08/16/89	07/12/89 08/15/89	
	Ship-		83	8	10	3	84	87	87	87	83	65	84	89 99	87	87	84	65	95	26	26	%	26	26	25	22	57	25	57	× 2	6	57	22	25	57	
	AVS		5058	5058	2058	2058	2905	2067	2905	5070	5070	2070	2205	202 202	8038	502	5073	2075	5098	2008	5113	5113	5133	5133	5134	5134	5137	5137	5138	5138	000	5142 5142	5153	\$153	\$156 \$156	

Ship- Test Plt ment Date # Diff. IC 25 TC 25 AI 25 IC 50 TC 50 AI 50 IC 95 TC 95 AI 95 SI TAI T se 07/25/89 QW1 1.341 13.20 0.96 0.07 23.60 64.90 2.74 0.00 268.00 0.00 0.04 5.46 58 07/25/89 QW7 1.180 88.50 196.00 2.21 0.00 296.00 0.00 0.00 320.00 0.00 0.00 5.18																	
Ship- Test Plt ment Date # Diff. IC 25 TC 25 AI 25 IC 50 TC 50 AI 50 IC 95 TC 95 AI 95 SI TAI se 07/25/89 QW1 1.341 13.20 0.96 0.07 23.60 64.90 2.74 0.00 268.00 0.00 0.04 5.46 se 10/04/89 RNV 0.852 0.00 16.00 0.00 0.00 49.00 0.00 0.00 301.00 0.00 0.00 0.00 se 07/25/89 QW7 1.180 88.50 196.00 2.21 0.00 296.00 0.00 0.00 320.00 0.00 0.00 5.18	4 ∪ ⊢	• •		+ + +	• •	+ +	+ +	• •	• • •	٠.	٠.	+ +	+ +	+ +	• •	٠.	
Ship- Test Plt ment Date # Diff. IC 25 TC 25 AI 25 IC 50 TC 50 AI 50 IC 95 TC 95 AI 95 SI T se 07/25/89 QW1 1.341 13.20 0.96 0.07 23.60 64.90 2.74 0.00 268.00 0.00 0.04 58 07/25/89 QW7 1.180 88.50 196.00 2.21 0.00 296.00 0.00 0.00 0.00 0.00 58 07/25/89 QW7 1.180 88.50 196.00 2.21 0.00 296.00 0.00 0.00 0.00 0.00	Assay Type	FF		THE THE	FF	T T T	# #	FF		E E	T I I	FE	FE	ĒĒ	EE	FE	
Ship- Test Plt ment Date # Diff. IC 25 TC 25 AI 25 IC 50 TC 50 AI 50 IC 95 TC 95 AI 95 58 07/25/89 QW1 1.341 13.20 0.96 0.07 23.60 64.90 2.74 0.00 268.00 0.00 58 07/25/89 QW7 1.180 88.50 196.00 2.21 0.00 296.00 0.00 0.00 320.00 0.00	Ĭ	5.46	5.18 27.60 1.30	14.49 7.73 0.00	2.46	22.86	10.88 23.33	8.04	60.23 5.80 13.82	5.34	13.86	23.45	31.32	49.55	29.05	69.04	
Ship- Test Plt ment Date # Diff. IC 25 TC 25 AI 25 IC 50 TC 50 AI 50 IC 95 TC 95 A se 07/25/89 QW1 1.341 13.20 0.96 0.07 23.60 64.90 2.74 0.00 268.00 se 07/25/89 QW7 1.180 88.50 196.00 2.21 0.00 296.00 0.00 0.00 320.00	3	0.00	8.30 0.00 v	0.00	0.00	2.64 >	1.91 > 2.64 >	88.50 v 0.00 v	320.00 > 0.00 > 2.24 >	32.70 > 0.00 >	13.10 > 0.00	4.81 > 5.51 >	6.95 >	75.20 > 159.87 >	38.20 >	217.00 > 0.00	
Ship- Test Plt ment Date # Diff. IC 25 TC 25 AI 25 IC 50 TC 50 AI 50 IC 95 58 07/25/89 QW1 1.341 13.20 0.96 0.07 23.60 64.90 2.74 0.00 58 10/04/89 RW 0.852 0.00 16.00 0.00 0.00 296.00 0.00 0.00 58 07/25/89 QW7 1.180 88.50 196.00 2.21 0.00 296.00 0.00 0.00 >	A1 95	0.0	0000	0.00	9.0	1.10 >	1.07 >	.00.0 .00.0	3.69 0.00	0.00	0.00	11.40 > 0.00	3.54	0.00	88.	37.40 > 0.00	
Ship- Test Plt ment Date # Diff. IC 25 IC 25 AI 25 IC 50 IC 50 AI 50 IC 95 58 07/25/89 QW1 1.341 13.20 0.96 0.07 23.60 64.90 2.74 0.00 58 07/25/89 QW7 1.180 88.50 196.00 2.21 0.00 296.00 0.00 0.00 58 07/25/89 QW7 1.180 88.50 196.00 2.21 0.00 296.00 0.00	10 95	301.00	320.00 1000.00 1000.00	307.00 95.60 293.00	257.00	320.00 >	320.00 >	320.00 320.00	320.00 × 305.00 320.00	96.40 29.40	90.60 77.30	29.80	320.00	320.00	320.00 320.00	320.00 > 320.00	
Ship- Test Plt ment Date # Diff. IC 25 TC 25 AI 25 IC 50 TC 50 58 07/25/89 QW1 1.341 13.20 0.96 0.07 23.60 64.90 58 07/25/89 QW7 1.180 88.50 196.00 2.21 0.00 296.00	10 95	0.0	0.00	9000	0.00	290.00 > 28.70	300.00 >	0.00	86.70 v 0.00 0.00 v	0.0	0.0	2.62	0.00 > 282.00 >	0.00	0.00	8.55 > 0.00 >	
Ship- Test Plt ment Date # Diff. IC 25 TC 25 AI 25 IC 50 S6 07/25/89 QW1 1.341 13.20 0.96 0.07 23.60 58 10/04/89 RWV 0.852 0.00 16.00 0.00 0.00 58 07/25/89 QW7 1.180 88.50 196.00 2.21 0.00	A1 50	2.74	0.00 17.76 0.00	0.00	2.01	2.64 61.73	1.91	196.00	320.00 0.00 2.24	55.50	25.20	8.63	6.95	75.20	84.60 0.00	217.00	
Ship- Test Plt ment Date # Diff. IC 25 TC 25 AI 25 IC 50 S6 07/25/89 QW1 1.341 13.20 0.96 0.07 23.60 58 10/04/89 RWV 0.852 0.00 16.00 0.00 0.00 58 07/25/89 QW7 1.180 88.50 196.00 2.21 0.00	TC 50	67.80	296.00 887.00 247.00	194.00 56.40 87.20	54.20 45.60	320.00 >	320.00 >	196.00 > 7.48	320.00 v 165.00 320.00 v	55.50 × 9.12	25.20 >	8.63 >	320.00 >	320.00 >	251.00	320.00 >	
Ship- Test Plt ment Date # Diff. IC 25 TC 25 15 56 07/25/89 QW1 1.341 13.20 0.96 58 10/04/89 RNV 0.852 0.00 16.00 58 07/25/89 QW7 1.180 88.50 196.00	22	23.60	00.00	0.00 23.70 0.00	27.00	121.00 >	168.00 > 379.00 >	0.0	1.00 > 0.00	0.00	0.00	1.00	49.00 ×	4.25 >	2.97	1.47 > 0.00 >	
Ship- Test Plt ment Date # Diff. IC 25 TC 58 07/25/89 QW1 1.341 13.20 0.58 10/04/89 RWV 0.852 0.00 16.58 07/25/89 QW7 1.180 88.50 196.	A1 25	0.00	2.21 18.58 1.13	8.85 2.41 0.00	1.46	9.31	2.6 10.00	88.50 × 0.00	320.00 < 0.00 6.48	32.70 < 0.00	13.10 < 0.00	4.81 <	32.00	232.00	71.00	320.00	
Ship- Test Plt ment Date # Diff. 58 07/25/89 QW1 1.341 58 07/25/89 QW7 1.180		16.00	196.00 414.00 90.30	130.00 34.60 55.10	19.40	320.00 >	320.00 >	88.50 > 2.59	320.00 > 90.60 320.00 >	32.70 >	13.10 >	4.81 > 7.43	320.00 > 715.00	320.00 >	114.00 38.80	320.00 > 320.00	
Ship- Test Plt ment Date # 58 07/25/89 QU1 58 07/25/89 QU1 56 07/25/89 QU7		13.20	88.50 22.30 80.20	14.30	13.30	34.40 >	121.00 >	0.00	1.00	0.00	0.00	1.00	10.00 >	1.38 >	1.60	1.00 •	
Ship- Test ment Date 58 07/25/89 58 07/25/89	Diff.	1.341	1.180 0.694 0.892	0.699	1.262	1.1 2 0.527	1.035	0.232 <	0.179 < 1.262 1.053	0.179 <	0.185 <	0.644 < 0.625	0.979	0.753	0.627	0.285 <	
S8 58 55 55 55 55 55 55 55 55 55 55 55 55	¥		PWZ VNB	RX0 ZAE QXE	02L S32	02P SY0	020 101	25	70 P 20	PCE PCE	P20	P3U SX2	P71 SY3	P7L SYO	P70 17X	100 TO	
		07/25/89 10/04/89	07/25/89 10/04/89 04/26/90	10/04/89 10/04/90 07/26/89	08/01/89 10/17/89	03/08/89	03/08/89 12/20/89	03/14/89 04/20/89	03/14/89 04/20/89 01/10/90	03/14/89 04/20/89	03/14/89	03/15/89	03/21/89	03/21/89	03/21/89 01/09/90	03/21/89	
AVS No. 174 174	Ship- ment	88	888	88	28 88	25	52	25	222	52	52	25	25	25	52	52	
		5174 5174	5186 5186 5186	5197 5197 5197	5210 5210	5241	5242 5242	5247	\$250 \$250 \$250	5251 5251	5252 5252	5253 5253	5271 5271	\$277 \$277	5283 5283	5291 5291	

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- 3	Assay Type	E E	ĒĒ	ĒĒ	EE	EE	E E	EEE	## ##					
	. Ε	14.28	0.74	16.05	27.11	21.97 6.88	0.00	2.73	11.87 7.98	24.31 0.00 3.51	10.98 0.00 5.04	2.54 12.47 1.65	23.12 22.95 7.26 0.00	22.90 20.25 22.17 8.06
	IS	2.15 > 1	0.00	2.83 > 1	18.50 > 2	1.20 > 2	1.18	0.00	2.18 1	6.33 × 2 0.00 0.00	2.16 > 1 0.00 1.31 >	0.00	5.16 × 2.37 × 0.00	3.62 × 2.56 × 2.
	AI 95	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00 0.00 0.00 0.00	0.00
	26 21	210.00	320.00 320.00	320.00 320.00	320.00 100.00	320.00 320.00	305.00	95.10 95.10 94.80	299.00	320.00 100.00 100.00	962.00 1000.00 320.00	32.00 320.00 100.00	320.00 > 1 32.00 100.00 94.50	320.00 966.00 955.00 320.00
	10 %	0.00	0.00.0	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00.0	0.00	2.83 0.00 0.00 0.00	00.00
nt'd)	AI 50	3.52	0.00	3.95	35.20	2.63	2.14 0.00	1.83	3.23	0.00	3.13 0.00 1.31	1.97	7.28 2.37 0.00 0.00	3.62 4.46 4.31 1.51
Table 21 (Cont'd)	10 50	61.70	25.30	320.00 >	320.00 > 70.50	180.00 67.30	169.00	50.70 51.40 47.70	91.60	29.30 25.90 18.30	618.00 1000.00 320.00 >	17.30 320.00 > 6.72	7.28 × 32.00 × 78.60 0.31	320.00 > 657.00 > 553.00 > 320.00 >
T.	10 50	17.50	0.00	81.10 × 0.00	9.09 × 7.69	00.0	0.00	27.70 0.00 0.00	28.40	2.81 0.00 0.00	198.00 0.00 > 245.00 >	8.79 179.00 > 0.00	13.50 0.00 0.00	88.40 > 147.00 128.00 > 211.00 >
	AI 25	6.36	1.38	6.00	168.00 132.00	38.80	1.86	1.62	4.59	14.20 0.00 2.50	3.08	3.01	5.16 × 6.03 3.26 0.00	7.82 7.50 12.48 3.03
	10 25	37.70	10.00	230.00	168.00 > 42.20 >	82.00	93.50	26.90 26.20 8.06	61.80	17.80 3.06 6.08	427.00 1000.0 320.00 >	320.00 >	5.16 × 32.00 × 55.30 × 0.32	320.00 × 485.00 329.00 × 320.00 ×
	10 25	5.93	7.26	38.30	1.00	2.11	50.30	16.70 18.30 0.00	13.50	1.25 0.00 2.43	139.00 0.00 > 156.00 >	5.14 106.00 > 2.83	5.31 × 17.00 × 0.00 ×	40.90 * 64.70 * 26.40 * 106.00 *
	01#	1.099	1.013	1.149	1.100 < 0.983 <	1.044	1.161 0.926	1.129 1.024 0.792	0.678	0.956	0.547 1.060 0.851	0.751 0.821 0.654	1.086 < 0.625 0.828 0.626	1.086 0.556 0.700 0.798
-	*	P18	PER	PVP	948	PWS 9A8	Pux 9A9	¥88	OAC OAC	PI7	SXV TOR	SXW WOF	916 8X2 8X1 XLT	710 VT1 V02
	Date	04/12/89	05/02/89	05/02/89 06/28/89	05/02/89	05/02/89 06/28/89	05/02/89 06/28/89	05/02/89 06/28/89 04/26/90	05/04/89 06/28/89	04/12/89 04/26/90 07/06/90	12/14/89 01/10/90 05/08/90	12/14/89 05/08/90 07/17/90	04/18/89 12/14/89 05/08/90 07/17/90	04/18/89 12/20/89 05/01/90 05/08/90
1:40	Bent .	88	22	22	22	22	22	228	22	55 53	223	233	8888	22.23
	. ÷	5350	5363	5367	5369	5372	5383	5384 5384 5384	2403	5450 5450 5450	2484 5484 5484	5485 5485 5485	2495 2495 5495 5495	5497 5497 5497 5497

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Account	Type	###				ĒĒ		ËË	ËË	Ħ	ĦĦ	ĒĒ	ĒĒ	ĒĒ	
	TA!	27.25 11.31 8.03	16.47 0.00 1.78	17.57 0.72 0.10	26.77 2.40 15.74	21.73	26.74 1.53 0.00 8.75	19.11	13.71	13.69	13.96	4.35	24.30	10.28	55.0
	2	2.2 2.35	2.47 > 0.00	3.19	3.92 > 0.00	2.27 > 0.00 >	2.38	1.30	0.00	2.49	3.05	3.70 >	5.10 >	1.51 >	0.00
	A1 95	0.00	0.00	999	0.00	0.00	00000	0.00	0.00	0.00	3.33	3.30	0.00	0.00	0.00
	tc 95	320.00 × 1000.00 320.00	9.72 320.00 10.00	304.00 300.00 262.00	320.00 309.00 307.00	328.00 97.40	198.00 259.00 86.40 312.00	320.00	320.00 942.00	320.00	320.00 320.00	320.00 966.00	320.00 954.00	320.00	265.00 75.30 32.00
	10 95	287.00 × 0.00 × 0.00 ×	0.00	0.00	00.00	0.0	0000	0.00	0.00	0.00	0.00 \$	293.00	0.00	0.00	0.00
nt'd)	AI 50	3.48	0.00	6.81 0.00 0.00	7.27 0.00 2.00	4.02	4.00 0.00 0.00 0.00	1.95	8.13	3.41	2.51	0.00	5.10	1.51	0.00
Table 21 (Cont'd)	16 50	320.00 × 689.00 320.00 ×	5.45 1.00 10.00	155.00 116.00 49.80	309.00 210.00 192.00	184.00 58.70	53.90 73.10 13.60 80.70	320.00 > 260.00	283.00	21.70	251.00 216.00	320.00	320.00 > 541.00	320.00 > 570.00	3.00
2	10 50	109.00 × 198.00 ×	0.00 0.00 0.00	22.80 0.00 0.00	42.50 0.00 96.00	00.0	0.00	164.00 >	0.00	6.37	100.00	0.00 >	62.80 > 72.80	213.00 > 233.00	0.00
	AI 25	3.58	3.08	8.44 0.00 0.00	9.33	6.52 3.72	7.02 0.00 0.00 3.76	7.38	5.71	4.26	3.64	1.82	7.61	2.58	3.30
	10 25	320.00 > 446.00 > 320.00 >	3.08	72.70 69.00 24.10	167.00 155.00 128.00	104.00 37.20	25.80 52.10 5.72 52.40	320.00 >	192.00	15.90	175.00 158.00	. 320.00 >	320.00 > 312.00	320.00 > 354.00	4.81
	10 25	48.40 > 125.00 140.00 >	0.00	8.62 0.00 0.00	17.90 0.00 34.70	15.90	3.67 0.00 0.00 13.90	43.40 > 16.50	33.60	3.72	48.20	176.00 × 35.90	42.00 > 37.50	124.00 > 137.00	0.00
	Diff.	1.058 0.523 0.759	1.057 < 0.759 0.829	0.979 0.677 0.738	0.956 0.556 0.755	0.899	0.905 0.877 0.884 0.988	1.099	1.227	1.294	1.220	1.447	1.257	1.315	1.339 0.864 0.988
-		SXX SXX	PL9 KOK KBB	4 865	102 E	PLD	PLG TFJ W2S ZYF	OCP R2H	0CS R21	0CT R41	0CV R42	00H	843	002 R44	0E0 R44 ZYF
	Date	04/18/89 12/14/89 05/08/90	04/18/89 05/08/90 07/03/90	04/18/89 02/13/90 05/08/90	04/18/89 12/20/89 05/08/90	04/18/89	04/18/89 01/18/90 05/10/90 11/01/90	06/13/89 08/08/89	06/13/89 08/08/89	06/13/89 08/09/89	06/13/89 08/09/89	06/14/89	06/14/89 08/09/89	06/14/89	06/14/89 08/09/89 11/01/90
Chio.	Ment	223	233	823	822	8 83	2822	92	28	22	26 56	28	28	28	288
AVC		2498 2498 5498	5500 5500 5500	5503 5503 5503	5507 5507 5507	5508 5508	5515 5515 5515 5155	5520 5520	5525 5525	5528 5528	5531 5531	5532	5534	5538 5538	5539 5539 5539

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	Assay Type	H H	FEEF	EE EE		FF	EEE	ĒĒ	ĒĒ	ĒĒ	ĒĒ	FFFF	ĒĒ	##
	IĀ	13.36	18.77 0.23 11.82	9.02	4.7.8	26.01 23.34	33.78 54.56 0.00	15.36	10. 87 2.91	9.02	10.84	4.78 5.98 0.33 9.61	2.30	3.12
	ïs	2.81 >	2.10 × 4.52 0.00 2.58 ×	0.00 > 5.92 > 313.00 >	17.74 > 19.17 > 0.01	4.94 >	6.69 × 24.77 × 0.00	2.18 > 0.00 >	1.60 \$	0.76 > 2.69	1.94 > 0.00	0.48 0.00 1.70	0.20 >	2.94 > 0.00 >
	AI 95	0.00	0.000	0.00	3.35	0.00	10.20 >	0.0	0.00	0.0	0.00	0.000	0.0	0.00
	TC 95	320.00	320.00 10.00 32.00 100.00	320.00	← W ← W	320.00 1000.00	320.00 320.00 > 307.00	320.00 100.00	92.60 96.10	320.00 940.00	320.00 939.00	9.54 5.14 9.60	320.00	320.00 320.00
	10 95	0.00	9888	0.00 0.00 789.00 4.00		0.00	0.00 × 31.40 × 0.00	0.00	0.00	0.00	0.00	8888	0.00	0.00
	AI 50	3.86	3.73 3.73 3.73	10.27	20.01	10.00	9.89	4.19	2.82	1.11	0.00	2.62 0.00 2.53	0.30	4.05
	Table 21 (Cont'd) TC 50 AI 5	7.06 10.00	17.80 1.81 32.00 26.20	173.00	1000.00 3 3200.00 3 686.00	320.00 > 1000.00 >	283.00 320.00 > 69.80	129.00	26.50 61.40	320.00 >	320.00 > 393.00	5.41 2.17 2.19 6.05	30.30	222.00
1	10 50	1.83	4.53 0.23 7.01	0.00 16.90 320.00 ×	50.00 × 128.00 × 246.00 104.00	64.80 v	28.60 12.90 v 0.00	30.90	9.39	288.00 >	141.00 >	2.06	100.00	0.00
	AI 25	4.81 0.00	5.08 7.34 0.00 4.11	0.00 24.88 313.00 <	156.74 48.05 0.02 31.56	28.97	26.50 47.84 0.17	5.27	3.54	6.83	5.15	< 0.93 2.16 0.00 3.12	0.40	4.32
	TC 25	5.13	9.50 1.06 25.90 18.10	100.00		320.00 3	192.00 320.00 3	67.40 25.10	15.00	218.00	274.00	1.57	19.70	161.00
	10 25	1.07	1.87	0.00 4.02 4.320.00 >	5.66 50.90 119.00 38.90	32.90 >	7.22 6.69 > 44.30	12.80	4.25	32.00 13.00	53.10	0.73 1.31	49.10	37.30
	oiff.	1.113	1.231 0.846 0.743 0.762	1.130	0.664 0.633 1.049 0.894	0.991	1.215 0.690 0.728	1.104	1.104	1.173	1.002	1.223 0.943 0.971 0.762	0.548	0.576
	# *	9E1	062 R46 VT2 ZAF	R47 PYG PYG		OLE R48	A 7 2 V T.3	ROE ROE	R9F	SZ SZ	R7	020 RVM S33 2AG	RPK 114	RPL T15
	Test	06/14/89 08/09/89 6	06/14/89 08/09/89 05/01/90 10/04/90	08/09/89		06/28/89	07/06/89 08/16/89 05/01/90	07/06/89 08/22/89	07/06/89	07/18/89 08/16/89	07/18/89 08/16/89	08/01/89 10/03/89 10/17/89 10/04/90	10/10/89 01/23/90	10/10/89 01/23/90
	Ship- ment	25.25	8888	22 22	62 GABSN 67 62/67	57	57 29	57	57	57	57	8 8 8 8	59	26
	AVS S	5542 5	5543 5543 5543 5543 5543	5548 5 5548 5 5580 5		5625 5	5643 5	5652 5	5653 5	5678 5678	5693	5714 5714 5714 5714 5714	5774 5	5780 5780 5
	~ ~	5 5	22.22.22	22 22 22	2222	2 2	222	2 2	2 2	2 2	2 2	2222	2 2	20.00

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	Assay Type	ĒĒ	ĔĒ	EE	ĒĒ	ĦĦ	# #	FF	ĒĒ		ĒĒ		ĒĒ	ĒĒ	ĒĒ		EE	
	IĀI	18.50	4.97	15.9	27.06	5.95	11.35	7.66	12.40	7.02 25.57 7.55	1.55	13.08 18.89 22.90	11.00	13.78	24.85	6.35	20.87	
	5	2.40 >	1.27	3.41	8.06 0.00	2.72 > 0.00	1.62 > 0.00	0.00	2.40 > 1.97	2.29 > 2.51 > 0.00	0.00	1.97 × 3.14 4.09	0.00 >	1.79 > 36.32 >	2.35 >	0.00 > 2.74	1.79 •	
	A1 95	88	0.00	0.00	0.00	0.00	0.00	9.0	0.00	0.00 0.00 0.00	0.00	1.07 × 3.24 3.35	0.00	1.06 >	3.25	3.20	0.00	
	70 %	320.00	320.00 320.00	311.00	30.30	9.51	98.20 95.90	320.00	320.00	320.00 32.00 32.00	306.00	320.00 > 966.00 963.00	320.00 320.00	320.00 > 320.00	320.00 > 966.00	320.00	320.00 > 979.00	
	1C 95	8.8	0.00	88	3.20	88	0.00	0.00	. 68	80.00	88	299.00 × 298.00 × 287.00	0.00	302.00 >	294.00 >	302.00	302.00 > 0.00	
(p,1)	A1 50	3.28	2.29	6.24	15.08	10.45	0.00	1.25	9.84 3.98	8.40 8.67 0.00	3.18	1.97 2 4.23 2 5.78 2	0.00 32.00	1.79 3	2.35 2 4.31 2	3.69	4.00	
Table 21 (Cont'd)	TC 50	216.00 62.60	216.00	171.00	15.10 > 3.20	10.40 > 5.07	57.40 59.00	320.00 >	28.30	8.40 > 32.00 > 29.60	183.00 209.00	320.00 × 660.00 630.00	320.00 320.00 >	320.00 >	320.00 >	320.00 >	320.00 >	
- T	10 50	0.00	94.00	27.50	0.00	0.0	21.40	256.00 > 0.00 > 1	2.68	3.69 >	0.00	163.00 × 156.00	0.00 •	179.00 × 8.81 ×	136.00 >	241.00 >	179.00 >	
	AI 25	0.00	2.74	7.32 2.14	8.06 <	2.72 <	2.7 0.00	3.56	3.06	2.29 < 10.28 5.60	1.92	2.76 4.50 7.50	0.00	2.39	6.22	3.66	17.89	
	TC 25	158.00	119.00	93.50 59.20	8.06 > 2.10	2.72 > 0.87	34.70	257.00	6.91	2.29 × 9.27 14.80	114.00 153.00	320.00 > 490.00 446.00	320.00	320.00 > 320.00 > 3	320.00 > 490.00	0.73	320.00 > 2.76	
	10 25	0.00	43.50	12.80 27.70	1.00	1.00	12.60	72.20	1.49	1.00 2.80 2.64	59.30	116.00 × 109.00 59.40	0.00 >	134.00 > 1.00 > 1.00	51.40 > 1	155.00	17.90 > 181.80	
	Diff.	0.547	0.805	0.860	0.821 < 0.885	0.821 < 0.763	0.933	0.838	0.918	0.918 < 0.632 0.553	0.710	0.638 0.778 0.583	0.623	0.663	0.663	0.594	1.024	
	Test Pit Date #	10/10/89 RPH 01/30/90 THL	01/30/90 TMM 01/31/90 TQM	10/31/89 SB1 01/30/90 THN	10/31/89 SB8 01/30/90 TMO	10/31/89 SBB 01/30/90 TMP	10/24/89 S6R 01/25/90 TJO	11/07/89 SF3 04/18/90 VCL	11/07/89 SF5 11/01/90 ZYH	11/07/89 SF5 02/01/90 TOE 02/08/90 TVX	11/09/89 SH7 02/01/90 TOF	01/31/90 TOP 10/30/90 ZW6 12/05/90 10F	01/31/90 TOR 12/05/90 10F	01/31/90 TOS 12/05/90 10G	01/31/90 TGS 10/30/90 ZW7	01/31/90 TQV 10/30/90 ZV9	01/16/90 TD7 11/01/90 ZYH	
	Ship-	59 1	29 0	2.0	19	10	33	1 19	1 19	199	1 0	62 0	62 0	62 0	62 0	62 0	62 0	
	No.	5782	5818	5879	5905	2906	5936 6	2004	5997 5997	5998 5998 5998	6209	6195 6195 6195	6200	6201	6202	6207	6214 6214 6214	

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	Assay Type	FF	FF	FF	FF	EEEE	FFF	EEE	FF	FF	FE	FF	FF	ĔĔ	FE	EE	
	TAI	13.74 24.28	14.47	1.83	16.79	17.58 30.51 6.74 1.30	1.34 24.69 6.28	0.30	7.87	24.26	0.00	8.26	39.78	4.89	0.40	3.09	
	25	1.66	1.97 >	0.00	2.19 > 3.40 >	0.00 0.00 0.00 v	0.00 5.85 v 0.00	0.00	0.00 >	2.87 > 0.00	1.79	0.00	8.14 >	1.79	0.00	0.42	
	A1 %	0.00	1.07 >	0.00	1.08 >	00000	0.00	0.00	0.00	0.00	0.00	0.00	3.51 >	0.00	0.00	0.00	
	8	320.00	320.00 >	320.00	320.00 > 966.00	1000.00 3200.00 3200.00 3200.00	1000.00 3200.00 3200.00	29.40 30.80 30.90	98.50 31.10	320.00 932.00	94.80	137.00	320.00 >	320.00 959.00	137.00	251.00	
	S 21	0.00	299.00 > 286.00 >	0.00	296.00 > 295.00	00000	0000	0.00 0.00 3.16	0.00	0.00	0.00	0.00	91.10 >	0.00	0.00	0.00	
nt'd)	A1 50	3.20 15.58	3.06	1.02	2.19	0.00 0.00 0.00	5.85	0.00	0.00	0.00	0.00	2.13	8.14	0.00	0.00	2.32	
Table 21 (Cont'd)	10 50	320.00 >	320.00 > 320.00 >	320.00 > 430.00	320.00 >	1000.00 3200.00 > 2780.00 3200.00	1000.00 3200.00 > 3200.00	9.06 19.60 20.70	63.70 21.10	320.00 > 320.00	48.50	\$1.00 67.70	320.00 >	320.00 585.00	43.50	55.50 83.50	
<u>1</u>	10 50	100.00 > 20.50 >	163.00 >	313.00 > 0.00	146.00 >	48.00.00 0.00.00	0.00 >	0.00	0.00	78.50 > 0.00	32.00	23.90	39.30 > 12.30	0.00 > 211.00	0.00	23.90	
	A1 25	8.44	2.76 5.73	1.48	3.40	312.50 10.67 3.13 0.00	0.84 10.46 3.12	0.00	6.49	28.26	0.11 25.64	3.83	23.56	1.50	0.90	0.86	
	10 25	166.00	320.00 >	262.00	320.00 >	1000.0 > 3200.0 > 1610.0 3200.0	773.00 3200.0 > 2100.0	6.13 13.50 15.00	15.60	225.00 116.00	2.67 57.40	18.50	320.00 > 613.00	249.00 378.00	8.23 0.58	10.00	
	10 25	12.90	116.00 >	0.00	94.20 × 86.70	300.00 × 513.00 × 0.00 ×	919.00 306.00 > 674.00	0.00	6.84 0.94	7.96	23.60	4.82	13.60 >	166.00 122.00	9.16 6.99	11.60	
	Diff.	0.831	0.943	0.943	0.908	0.619 0.586 0.902 0.759	0.604 0.548 0.888	0.539	0.605	0.449	0.940	0.823	0.724	0.773	0.750	0.782	
	÷ *	109 10H	10A 2Y1	101 101	100 271	SS1 TW7 W14 ZYJ	SSJ TWB	UZK UPO ZAH	USN UNR	U8G UPV	W2Z XDG	X 0 X	¥32	No.	15 V	W57 XF0	
	Test	01/16/90	01/16/90	01/16/90	01/16/90	12/05/89 02/08/90 05/31/90 11/01/90	12/05/89 02/08/90 05/31/90	02/15/90 03/20/90 10/04/90	02/15/90	02/27/90	05/10/90 07/06/90	05/10/90 07/06/90	05/10/90 07/06/90	05/15/90 07/06/90	05/15/90 07/06/90	05/15/90 07/10/90	
	Ship-	33	33	33	2 29	62 GABSN 67 67	62 GABSIN 67	222	នួន	នន	88	88	88	33	33	88	
	No.	6218 6218	6219	6220	6225 6225	6234 6234 6234	6236 6236 6236	6315 6315 6315	6321	6373 6373	6412	6413	417	6422	6435	441	

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	Type	FE	FE	<u> </u>	FE	FF		HH	FF	TH	FE	Ħ Ħ	ĦĦ	FF	H	FE	EE
	TAI	13.95	18.41	5.69	11.40	9.84	21.03	12.73 25.64	6.97	11.10	3.8	14.97	4.81	5.85	0.00	0.00	6.49
	Is	1.85 v	4.82	1.24	99.9	1.50 > 22.43 >	0.00 6.35 2.33	3.89	0.00 >	3.16	0.00	2.56	. 200	0.00	0.00	0.00	0.00
	VI 88	3.39	% % %	9.0	88	0.00 >	888	88.	9.5	3.41	88.	88	88	88	88	88	8.8
	TC 95	307.00	320.00	95.20 96.60	320.00	320.00	320.00 320.00 100.00	320.00 1000.00	320.00	310.00	9.61	96.60	30.60	30.80	137.00	320.00	320.00 954.00
	10 95	0.00	0.00	0.00	0.00	400.60	0000	0.00	0.00 >	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00
nt'd)	AI 50	2.86	0.00	2.23	0.00	1.50	0.00 25.15 5.67	3.89	0.00	5.43	0.00	3.45	0.00	2.17	0.00	0.00	0.00
Table 21 (Cont'd)	10 50	187.00	320.00 >	52.10 66.00	87.70	320.00 >	24.70 73.80 40.50	320.00 >	320.00	160.00	6.09	21.50	18.30	3.01	43.50	320.00	295.00
7	10 50	65.40	66.40 00.00	23.40	0.00	213.00 > 44.60 >	0.00 2.94 7.14	82.30 > 86.20 >	73.10	65.80 33.40	0.00	19.20	0.00	9.19	9.70	0.00 >	0.00
	AI 25	3.59	6.94	1.97	7.87	3.11	1.72 10.88 3.49	8.78	1.98	3.65	2.28	3.92	2.67	2.56	0.98	0.00	3.26
	TC 25	121.00	320.00 > 384.00	29.00	59.90	320.00 > 1000.0 >	6.71 18.60 16.70	320.00 > 759.00	100.00	85.10 106.00	4.14	49.00	11.40	13.90	14.40	320.00	189.00
	10 25	33.70	46.10 > 0.00	14.70	3.98	103.00 >	3.90 1.71 4.78	36.50 ×	50.50	23.30	1.81	12.50	4.25	5.42	14.70	0.00 >	57.90 23.50
	Diff.	0.825	0.752	0.919	0.888	0.742	0.828	0.781	0.770	0.749	0.623	0.623	0.729	0.761	0.747	0.732	0.732
2	: *	XFO X	VEN	V40 X70	VIS	KK2 XHF	XXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXXX	XX	Z X	₹ ×	X X	XLN	XNS	XNU	E K	XRT	TNS T
	Date	05/17/90 1	05/30/90	04/02/90	05/31/90	06/05/90	06/05/90 07/12/90 11/15/90	06/05/90	06/01/70	06/12/90	04/11/90	07/17/90	06/119/90	07/19/90 XNU 09/05/90 YKY	04/19/90	04/54/90	04/56/90
	ment ment	33	88	33	79	79	55 57	67	79	22	69	88	69	69	69	69	\$ \$
	NO.	6482	6570 6	85.78 87.89	6 7076 6 7076	6714 6	6724 6724 6724 674	6753 6	6788 6 6788 6	6837 6	9 9889	6 7889 6 7889	9 9689	9 0069	9 9069	6942 6	6943 6
•	. =	22	25	33	67	67	67	55	67	28	28 28	2 2	28.28	59 59	500	\$ \$	69 69

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	od.	EE	T I I	FE	EEE	FFF	EEE	FF	FER	E K	T T T T	FF	# E	EE	FEE	EE	
	TAI	10.26	6.7	0.40	2.47 8.00 1.66	8.88 13.98 19.75	21.98 32.48 14.74	11.45	13.46	25.88	17.80	0.00	3.69	36.89	18.27	0.83	
	75	1.85	0.0	0.00 > 2.46	0.00	1.44 2.87 2.48	3.53 × 5.07 × 1.73 ×	1.71 >	1.92 > 3.16	5.19	3.89 >	1.86 \$ 0.00	1.32	0.03 >	2.68 > 0.00	0.00	
	A1 95	0.00	0.00	0.00	0000	3.39	3.20	0.00	3.07	3.60	0.0	1.06 •	0.0	0.00	1.10	00.0	
	10 95	320.00	320.00	320.00	305.00 307.00 100.00	306.00 309.00 308.00	309.00 309.00 306.00	320.00	320.00 966.00	320.00 > 963.00	320.00 993.00	320.00 > 313.00	306.00	88.30 1000.00	320.00 > 294.00	320.00	
	10 %	0.00	0.00	0.00	0.00	0.00 0.00 0.60	96.60 89.00 0.00	0.00	315.00	88.80 × 88.10	0.00	301.00 \$	0.00	302.00	290.00	0.00	
ont'd)	A1 50	3.81	0.00	0.00	0.00 0.00	3.90	6.88 2.95	1.71	1.92	7.15 22.44	3.07	1.86	2.21	0.05	2.68	0.00	
Table 21 (Cont'd)	10 50	223.00	316.00	320.00 558.00	165.00 186.00 67.80	179.00 209.00 168.00	210.00 210.00 175.00	320.00 >	320.00 >	216.00 634.00	320.00 > 674.00	320.00 >	177.00	8.68	320.00 >	320.00	
2	10 50	54.00 61.60	95.00	0.00 > 137.00	0.00	75.20 53.50 37.40	43.90 30.50 59.30	187.00 > 0.00 >	167.00 >	30.20	71.40 > 220.00	172.00 >	80.20	179.00	119.00 >	312.00	
	A1 25	4.36 3.50	8.33	0.00	1.58 2.51 0.00	3.18 5.38 5.02	6.80 16.81 5.50	2.34	4.14	10.74	5.80	2.53	2.09	38.73	0.00	0.00	
	7C 23	100.00 96.00	208.00	338.00	90.60 119.00 49.90	108.00 153.00 92.70	155.00 155.00 103.00	320.00 >	320.00 >	157.00	277.00	320.00 >	106.00	5.94	320.00 >	228.00	
	10 25	23.00	24.90	0.00	\$7.20 47.30 0.00	34.10 28.50 18.50	22.80 9.22 18.70	107.00 >	7.30 >	14.60	47.80	127.00 >	50.70	134.00	0.00	0.00	
	oiff.	0.796	0.546	0.789	0.816 0.683 0.709	0.909	0.909	0.856	0.856	0.849	0.764	0.716	0.572	0.749	0.617	0.867	
-	ž *	XAU	TNT	XUX	XXX YTN	XZZ XXY YTN	XXX 410	X23	X23	X27	X28	YNC	XUX	3 S	XWS TOT	35	
	Oate	07/24/90	07/26/90	07/26/90	06/21/90 08/02/90 09/13/90	06/21/90 08/02/90 09/13/90	06/21/90 08/02/90 09/13/90	06/21/90	06/21/90	06/21/90	06/21/90	07/26/90	07/26/90	07/31/90	07/31/90	07/31/90	
	ment ment	99	\$ \$	66	333	333	333	33	33	2 2	22	99	99	6 6	99	60	
	N 0.	7769	6945 6	9769	0269	9769 9769 9769	6977 6977 6977 6977	87.69	6269	9869	9869	7003	7022	7023 6	7032 6	7036	
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Assay Type	EE	##	FF	FE	FF	EEE	EEE	EEE		EEE	EEE	EEE	
IAI	5.81 3.59	15.92	5.22	0.00	15.67	7.05 0.00 6.79	90.09	1.82 3.11 10.15	6.00.4 0.00.8 0.00.8 0.00.8	8.91 1.88 12.51	6.11 0.00 15.48	3.39	35.95 16.41 12.33
3	0.00	2.62	0.00	2.52 > 0.00	2.19 >	1.67	1.07 > 0.00	0.00	0.0000000000000000000000000000000000000	1.24 > 0.00	0.00	0.00	10.22 > 3.15 > 2.99
A1 95	0.00	3.28	0.00	0.0	0.00	0.00	0.00	0.00	, 9000000000000000000000000000000000000	0.00	0.00 0.00 3.16	0.00 0.00 v	0.00
25 25	320.00	309.00	307.00	308.00	320.00 >	309.00 296.00 307.00	100.00 691.00 320.00	320.00 955.00 1000.00	309.00 300.00 100.00 932.00	320.00 942.00 963.00	320.00 935.00 986.00	320.00 1000.00 1000.00	304.00 297.00 299.00
1C 95	0.00	94.20	0.0	0.0	296.00 > 0.00 >	988	0.00	988	8000000	0.00	0.00 0.00 312.00	0.00	31.10 0.00 0.00
AI 50	0.00	3.83	2.02	3.54	3.11	2.31	1.07	0.00	2001.00 2000.00 2000.00	1.37 0.00 2.63	0.00	0.00	18.90 4.90 6.71
Table 21 (Cont'd) 1C 50 A1 5	320.00	210.00	194.00	203.00	320.00 > 699.00	207.00 86.90 189.00	100.00 × 145.00 215.00	320.00 547.00 602.00	208.00 48.70 124.00 100.00 v 320.00	320.00 > 424.00 630.00	320.00 346.00 664.00	320.00 > 233.00 427.00	161.00 97.20 112.00
Ta 10 50	0.00 > 259.00	54.90	%. 0.00	0.00	146.00 > 225.00	89.60 0.00 68.40	93.40 × 0.00 85.00	0.00 > 184.00	20.00 0.00 0.00 0.00 0.00	234.00 > 0.00 239.00	0.00 > 182.00	292.00 > 0.00 0.00	8.50 19.80 16.80
S 14	3.20	3.81	2.38	3.59	3.34	2.89	1.89	0.00 1.66 6.16	4.74 0.00 1.81 0.00 0.45	3.64 1.13 3.26	3.59	4.00 0.00	20.14 4.84 6.06
TC 25	320.00 > 231.00	155.00	130.00	144.00 34.40	320.00 >	150.00 28.20 124.00	100.00 × 43.30 128.00	320.00 320.00 341.00	152.00 25.60 38.50 100.00 • 143.00	290.00 230.00 446.00	283.00 162.00 485.00	320.00 > 90.30 191.00	86.80 62.50 50.10
52 21	100.00 >	0.00	54.70	40.20	95.70 ×	51.80 0.00 46.80	52.80 > 0.00 47.50	0.00 × 193.00 55.40	32.00 0.00 0.00 55.20 >	79.60 204.00 137.00	158.00 0.00 135.00	165.00 v 0.00 0.00	4.31 12.90 8.27
oiff.	0.655	0.500	0.734	0.776	0.756	0.823	0.655 0.785 0.681	0.730	0.721 0.808 0.739 0.739 0.706	0.763	0.747	0.747	0.718 0.863 0.601
<u>:</u> *	Z X	YRK	YRL	YRM	YRN	YE2 288 20U	YE3 28C 20V	YE4 280 29V	7FY 20Y 7G0 78F 20Y 20Y 20Y	YG0 28F 20Z	761 286 202	YG1 28G 2R0	766 28H 2R0
Test Oate	07/31/90	07/31/90	08/02/90	08/02/90 09/11/90	08/02/90 09/11/90	08/29/90 10/02/90 10/25/90	08/29/90 10/02/90 10/25/90	08/29/90 10/02/90 10/25/90	08/30/90 10/02/90 10/25/90 08/30/90 10/02/90	08/30/90 10/02/90 10/25/90	08/30/90 10/02/90 10/25/90	08/30/90 10/02/90 10/25/90	08/30/90 10/02/90 10/25/90
Ship-	\$ \$	69	69	69	69	222	222	222	22222	222	222	222	222
AVS.	7040	7042	300	7048	7049	7055 7055 7055	7057 7057	7059 7059 7059	7068 7068 7071 7071	2707 2707 2707	55 57 57 57 57	707 707 707	7083 7083

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	Assay	FFF			***	FFF	FE	H H	H H	E E	H	EE	H H	EE	HH	Ħ.H	
	TAI	38.97 16.60 17.48	30.82 14.51	21.69 8.81 26.98	35.17 23.48 22.80	15.65 22.55 32.11	0.00	7.45	6.58	4.37	3.25	0.00	9.43	11.31	3.56	11.52	
	22	12.21 > 2.30 > 1.65	6.44 2.83 v 0.22	4.9% > 1.97 7.87 >	4.04	1.96 > 2.38 > 0.03 >	0.00	1.88	1.09 •	1.09	1.10	1.55	1.41 > 0.00	1.00 \$	1.05 >	1.69 \$	
	A1 95	11.57 0.00 0.00	11.03	10.26 0.00 0.00	0.00 0.00	0.00	0.0	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	
	10 %	%.30 91.80 93.30	100.00 273.00 100.00	29.90 85.20 88.70	100.00	320.00 1000.00 >	320.00	311.00	1000.00 3200.00	313.00	305.00	100.00 30.70	320.00	1000.00 3200.00	320.00	320.00	
	10 %	8.32 0.00 0.00	9.00 0.00 v	0.00	9.30 \$	0.00 > 947.00 > 1	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00	
nt'd)	A1 50	21.78 4.16 22.35	12.11 5.19 11.42	7.32 3.02 13.73	31.98 6.03 16.53	1.96 2.38 10.14	0.00	2.62	0.00	2.08	3.01	3.23	1.41	1.30	1.05	1.64	
Table 21 (Cont'd)	10 50	40.70 28.00 33.30	45.40 40.70 42.10	9.16 23.30 23.90	31.10 29.30 23.30	320.00 × 1000.00 × 1000.00 ×	54.70	207.00 370.00	1000.00 > 3200.00	166.00	169.00	19.40	320.00 >	1000.00 >	3200.00	320.00 >	
T	10 50	1. 27 6.73 1.49	3.74 7.85 3.69	1.25	0.97 4.86 1.41	164.00 × 420.00 × 98.60 ×	0.00	0.00	918.00 × 0.00 ×	79.60	84.00 67.60	20.60	228.00 > 0.00	769.00 \$	305.00 >	174.00 > 0.00	
	AI 25	29.65 8.43 4.20	13.47 5.19 0.44	6.18 3.24 12.88	19.92 5.80 8.19	3.69 6.79 0.10	0.00	3.22	2.22	1.92	1.78	2.26	2.86	0.00	1.92	5.41	
	10 25	22.80 15.50 2.47	24.10 22.20 0.83	6.18 × 15.20 13.70	9.50 19.60 5.75	320.00 > 1000.0 > 2.67	10.00	149.00	1000.0 > 3200.0	86.40 146.00	92.40	32.00	320.00 > 287.00	766.00 3200.0	320.00 > 2320.0	294.00	
	10 25	0.77 1.84 0.59	1.79	1.06	3.39	86.70 > 147.00 > 26.30	0.00	46.10	450.00 >	45.00	51.90	14.20	112.00 >	179.00	166.00 \$	54.30	
	Diff.	0.863	0.79	0.674 < 0.799 0.799 0.716	0.688	1.050 0.708 0.569	0.771	0.763	0.735	0.710	0.725	0.716	0.565	0.660	1.001	0.705	
	= *	766 28H 2R1	767 281 281	767 281 282	768 28J 2R2	28 X X X X X X X X X X X X X X X X X X X	YOY	747	Y50	754 7TX	YSS	Y58	774 748	Y73	178 148		
	Test	08/30/90 10/02/90 10/25/90	08/30/90 10/02/90 10/25/90	08/30/90 10/02/90 10/25/90	08/30/90 10/02/90 10/25/90	09/05/90 10/02/90 10/25/90	08/07/90	08/14/90	08/14/90	08/14/90	08/14/90	08/14/90	08/16/90	08/16/90 Y79 09/18/90 WA	8/16/90 Y78 09/18/90 YW	08/21/90 Y93 09/18/90 YWC	
	Ship-	222	222	222	222	222	22	22	22	22	22	22	22	22	22	22	
	No.	7064	7065	7086 7086 7086	7087 7087	7092 7092 7092	270	7320	7323	7332	7333	77.	7356	7365	7369	22 22 22	

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	Type	FIE	FER	FIR	FE	TIM TIM	EE	FE	FER	FIN	FER	FER	EE	THE	EE	FE	H H	
	¥	6.00	31.26	6.59	12.25	3.50	2.0 2.4	11.23	11.56	5.95	17.22 7.95	14.86 8.76	20.72	14.95	10.52	20.53	3.01	
	5	2.52	6.91	1.81 >	1.97 >	1.28 > 0.00	0.80	% 0.00	1.49 > 2.13	6.57 >	3.46	2.82	5.16 v 1.18	3.23	1.68 >	2.24 >	0.00 >	
	A1 95	88	0.00	1.06 0.00 v	0.00	0.0 0.0 v	0.0	1.07 >	0.00	0.0	9.24	3.24	0.0	0.00	0.00	0.00	0.00	
	70 %	966.00 3200.00	320.00	320.00 > 955.00	320.00	320.00	320.00 575.00	320.00 > 937.00	320.00 958.00	971.00	27.90	9.63	30.70	313.00	320.00 1000.00	320.00 1000.00	304.00	
	10 %	88.	0.00	302.00 >	88	88.	% 88.	300.00	9.0	88.	3.02	8.0	8.6	88.	0.0	0.00	9.0	
4,9	AI 50	3.40	0.00	1.86	1.97	1.28	0.00	4.00	1.49 3.34	9.15	4.92	4.12	2.05	5.01	1.68	2.24	3.36	
Table 21 (Cant'd)	TC 50	660.00 3200.00	1000.00 >	320.00 >	320.00 >	320.00 >	320.00 >	320.00 >	320.00 >	607.00	7.86	6.14	18.70	83.50 50.10	320.00 >	320.00 > 849.00	156.00	
-	10 50	194.00	112.00 > 0.00 >	177.00 > 294.00	162.00 >	249.00 >	246.00 >	165.00 >	215.00 > 174.00	00.00	1.60	1.49	2.35	17.90	190.00 >	143.00 >	0.00	
	VI 25	5.14	45.52	2.27	3.63	2.16	2.07	2.69	2.76	0.17	3.78	4.14	9.34	3.80	4.50	5.57	6.00	
	TC 25	490.00	1000.0 >	320.00 >	320.00 >	320.00 > 241.00	197.00	320.00 >	320.00 >	9.38	5.53	4.20	12.10	57.80	320.00	320.00 > 377.00	79.90	
	10.25	9.50	22.00 >	131.00 >	88.10 × 0.00	148.00 >	95.20	119.00 > 228.00	116.00 × 69.80	54.50	1.12	1.02	1.30	12.30	71.00	57.50 > 47.10	0.00	
	Diff.	0.711	0.770	0.743	0.643	0.820	0.765	0.781	0.722	0.829	0.706	0.705	0.821	0.691	1.095	0.989	1.286	
		₹ 38	797 23K	230	YBR 23V	YDU 23V	Y0V 23W	70V	Y0Y 28A	7Y7 2R4	77E 2V2	YYE 2VZ	TYF ZVO	241	20E	202	2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2	
	Date	08/21/90 195 09/18/90 TWE	08/21/90	08/23/90	08/23/90	08/29/90	08/29/90	08/29/90	08/29/90 YDY 10/02/90 Z&A	09/20/90 YY7 10/25/90 2R4	09/20/90 YYE 10/30/90 ZVZ	09/20/90 YYE 10/30/90 ZVZ	09/20/90 YYF 10/30/90 240	09/20/90 YYG 10/30/90 ZW1	09/25/90	10/30/90	09/25/90 20N 10/30/90 2W4	
	Sent Pe	22	22	22	22	22	22	22	22	22	22	RR	RR	RR	22	RR	RR	
		23.28	7382	7418	7424	7427	74.29	7430	76.34	7439	7457	7458	7459	1972	6972	7872	787	
				-				-			-	-	-			_		

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	Type		FF	TH T		ĒĒ	H H	H H
	IAI	3.33	0.00	3.07	0.00	3.01	0.35	16.91
	75	0.00	5.33 >	0.00	3.08	1.00	0.00	3.50
	A1 %	9.0	0.00	88	88	90.0	0.0	3.35
	TC 95	320.00	320.00	320.00	320.00	320.00 955.00	955.00 3090.00	3090.00
	10 %	0.00	0.00	0.00	0.00	0.00	0.00	922.00
ont'd)	AI 50	1.24	0.00	0.00	6.0 7.	_	3.48	3.73
Table 21 (Cant'd)	10 50	210.00	3200.00	320.00	762.00	320.00 547.00	95.30 2100.00	2050.00
2	10 50	804.00 \$	0.00 > 544.00 >	0.00 > 2620.00 >	0.00 > 176.00	320.00	0.00	443.00
	AI 25	0.00	0.00	0.00	0.00	0.00	0.16	5.88
	TC 25	26.30	320.00	218.00	245.00	97.40	36.70	1550.00
	10 25	439.00 >	0.00 > 320.00 239.00 2900.0	0.00	0.00	179.00	224.00	327.00
	Diff.	1.286	0.883	0.833	0.723	0.804	0.719	0.395
=		252 252	2GB 123	2K1				N H
	Dete	09/25/90 20N 10/30/90 245	10/11/90 2GB 11/06/90 123	10/16/90 2K1 11/08/90 141	10/16/90 2K6 11/08/90 146	10/16/90 2K9 11/08/90 148	12/11/90 1SX 01/31/91 2ZL	12/13/90
C) 45	S S	RR	22	22	22	% K	KK	22
	9	74.88	300	300	7935	20,00	6762	8374

This value is a virus rating (VR) rather than a TAI. The VR is a measurement of selective antiviral activity that takes into account the degree of inhibition of virusinduced CPE and the degree of cytotoxicity produced by the test compound similar to TAI. TAI is more accurate with MTT measurements.

DIFRNTL = The differential is the difference in the cell control and the virus control optical densities.

(Viral) inhibitory concentration 25%, 50% and 95% = The drug concentration (µg/ml) that inhibited viral CPE by 25%, 50% or 95% calculated by using a regression analysis for semilog curve fitting. IC25,50,95

TC_{25,50,95} = (Cell) toxicity concentration 25%, 50% and 95% = The drug concentration (µg/ml) that reduced cell viability by 25%, 50% or 95%.

Antiviral Index = A single point ration of the antiviral and anticellular effect of the compound, calculated with 25%, 50% or 95% reduction values (calculated by dividing the TC25,50,95 by the IC25,50,95). AI25,50,95

Selectivity Index = A ratio calculated by dividing the TC₂₅ by the IC₃₀ (based upon 6 one-half-log₁₀ dilutions, μ g/ml, the maximum scale is 0-320). S

Total Antiviral Index = The area between the cytotoxicity and the antiviral curves (based upon a scale of 0-100%). TA

Activity = A "+" denotes a test that produced >25% reduction in CPE. A "-" denotes an inactive test (i.e. <25% reduction in CPE. ACT

4.1.8 Hantaan Virus (HTN)

Initially, we had problems adapting a method for evaluating compounds against the Hantaan Virus (See section 4.1.10 of Second Annual report dated December 15, 1987). A plaque-reduction method was developed but it was unsuitable for our large-scale screening program. We continued to test compounds versus HTN by plaque-reduction assay through April, 1988. At this time, testing of the antiviral compounds against the HTN virus was suspended in our laboratory to complete the development of the ELISA system (a less labor-intensive and suitable for large-scale screening assay) under the direction of Dr. John Huggins at USAMRIID, Ft. Detrick, Maryland. However, testing of antiviral agents against HTN has continued to be carried out at USAMRIID under the direction of Dr. Huggins and Dr. Z. Zhang.

With the plaque-reduction assay, one compound, AVS-3593 showed moderate antiviral activity against the virus with an ID₅₀ of 0.93 μ g/ml, a MTC of 1.0 μ g/ml and a TI of 1.1. Ribavirin was our positive control compound and yielded ID₅₀'s ranging from 1 - 17 μ g/ml and TI values that ranged from 6.3 - 57 (based on results of 12 assays). Table 22 shows a comparison of AVS-3593 and Ribavirin tested on February 17, 1988.

Table 22

Compounds Active Against Hantaan Virus^a

AVS No.	IC ₅₀ (µg/ml)	MTC <u>ug/ml</u>	TI
0001	15.9	100	6.3
3593	0.9	1.0	1.1

Plaque-reduction assays were done in C1008 cells.

IC₅₀ = The minimum drug concentration (μ g/ml) that inhibited the CPE by 50%, calculated by using a regression analysis program for semilog curve fitting.

MTC = Minimum cytotoxic drug concentration is the lowest drug concentration at which drug toxicity was observed.

TI = The Therapeutic Index of a compound was determined by dividing the MTC by the IC_{50} .

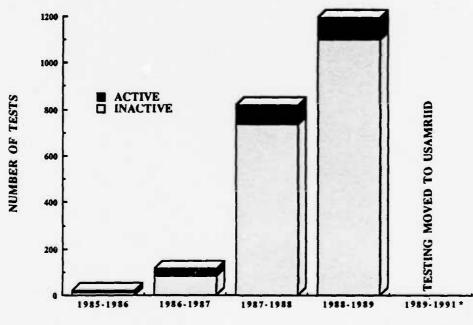
4.1.9 Pichinde Virus (PIC):

The number of single drug tests carried out against PIC during this contract period is summarized in yearly increments in Figure 42. During 1989, testing was moved to USAMRIID in order for SRI to develop an ELISA assay system suitable for primary screening of large numbers of compounds versus the Pichinde virus.

A total of 2276 tests were performed during this contract period using a plaque reduction assay procedure. Ribavirin (AVS-0001) was tested in each plaque reduction assay as the positive control compound. Data from 135 positive control assays tested during 1988 - 1989 yielded a mean IC₅₀ of 10.65 (\pm 5.4) and a range of 4.1 μ g/ml to 18 μ g/ml. The mean MTC was 76.8 μ g/ml (\pm 35.1). The mean TI value was 7.95 (\pm 4.33). Results of Ribavirin tested in parallel in each assay were used as a guideline to assess the quality of each assay in the day-to-day screening.

Out of the 2141 accepted single drug tests, 210 compounds demonstrated antiviral activity at greater than or equal to 50% reduction levels. This represents around 10% of the tested compounds having *in vitro* antiviral activity against the PIC virus. The remainder, 1931 compounds (90%), were considered inactive with the CPE-inhibition assay protocol (Figure 42).

IN VITRO PRIMARY SCREEN: NUMBER OF COMPOUNDS FOUND ACTIVE AGAINST PICHINDE VIRUS DURING THE CONTRACT PERIOD



Status	1985-1986	1984-1987	1987-1988	1988-1989	1989-1991-	Five-Year <u>Totals</u>
Number Active	9	22	83	96	0	210
Number Inactive	4	<u>92</u>	<u>736</u>	1099	Q	<u>1931</u>
Yearly Total (Accepted Single Drug Tests)	13	114	819	1195	0	2141

Represents 14-month period (November 15, 1989 - January 31, 1991)
Figure 42

4.1.9.1 PIC - Antiviral Activity Results:

Of the 2141 compounds tested against PIC using the plaque reduction assay, 197 (7%) demonstrated antiviral good activity with antiviral reduction values (IC₅₀) equal to or better than 50% and a Therapeutic Index of > 1.0 (Table 23). Five compounds (AVS-0646, 0148, 0140, 2350 and 3189) produced Therapeutic Indices (TI's) of > 100. Fifteen compounds produced good antiviral activity with TI's that ranged from > 10 - 62. These compounds should be tested further in *in vitro* profile studies and *in vivo* testing.

Table 23

Compounds Active Against Pichinde Virus^a

AVS No.	Shipment	<u>IC</u> 50	_MTC_	TI
0646	2	0.1	100	>681
0148	2	0.2	100	589
0140	4	0.5	> 100	> 208.2
2350	14	158	> 20000	> 126
3189	29	0.9	100	106
3800	35	0.16	10	62.4
0215	1	2.2	> 100	>45.4
0001	4	2.5	> 100	>40.1
2527	20	0.9	32	34
0230	1	9.8	320	32.7
0139	1	15.4	>320	>20.8
3799	35	0.56	10	17.6
3802	35	0.56	10	17.6
2581	19	5.9	100	17
0253	1	2.0	32	16
0272	1	2.0	32	16
2568	21	26	320	12
2585	19	8.3	100	12
0095	2	9.1	> 100	> 10.9
3801	35	0.31	3.2	10.1
4452	44	11	100	9.5 `
2506	21	1.4	10	7.4
1985	2	4.7	32	6.9
2503	21	0.14	1	6.9
2618	19	47	>320	>6.8
3803	35	0.49	3.2	6.5
2594	19	16	100	6.2
2979	25	17	100	6.0
2582	19	17	100	5.8
0332	4	17.5	> 100	>5.7
2591	19	17	100	5.7
0360	2	0.2	1	5.5
5480	53	60	320	5.3
4035	36	6.9	32	4.7
3038	28	0.06	0.32	4.7
4871	46	22	100	4.6

Table 23 (Cont'd)

AVS No.	Shipment	IC ₅₀	MTC	IL
2868	20	2.2	< 10	<4.5
3422	32	2.4	10	4.2
4240	39	2.5	10	4.0
0206	4	87.2	>320	>3.7
2577	19	28	100	3.6
3490	33	9.2	32	3.5
1992	3	94.9	320	3.4
0094	1	30	> 100	>3.3
2592	19	100	320	3.2
4768	44	31	100	3.2
4611	43	104	> 320	3.1
4234	42	35	100	2.9
3206	29	1.2	3.2	2.8
0068	1	112.3	>320	>2.8
1975	· i	120.6	> 320	>2.7
4073	37	119	320	2.7
1378	45	39	100	2.6
2576	19	38	100	2.6
2575	19	40	100	2.5
2125	7	13	32	2.5
0111	22	40	100	2.5
2584	19	42	> 100	
				>2.4
2980	25 24	1.4	3.2	2.4
2811	24	0.13	0.32	2.4
4434	44	13	32	2.4
4113	39	1.4	3.2	2.4
2089	6	4.3	10	2.3
2277	11	43	100	2.3
3190	29	14	32	2.3
4887	46	1.5	3.2	2.2
0113	1	45.1	100	2.2
0347	2	0.4	1	2.2
2132	7	45	100	2.2
2589	19	48	100	2.1
4610	43	155	320	2.1
4190	39	16	32	2.1
4464	45	0.16	0.32	2.0
2363	15	16	32	2.0
2172	8	16	32	2.0
3503	33	16	32	2.0
0708	30	50	100	2.0
3565	32	5.0	10	2.0
0349	2	1.6	3.2	2.0
2108	6	52	> 100	>1.9
0084	1	172	320	1.9
4071	35	53	100	1.9
4273	39	17	32	1.9

Table 23 (Cont'd)

AVS No.	Shipment	IC ₅₀	_MTC_	IL
4272	39	56	100	1.8
2630	19	57	100	1.8
2863	20	1.8	3.2	1.8
2907	26	57	100	1.7
3141	29	5.7	10	1.7
2228	10	19	32	1.7
3073	28	1.9	3.2	1.7
1970	27	19	32	1.7
4390	43	18	32	1.7
4769	44	19	32	1.7
5558	53	19	32	1.7
2408	16	192	320	1.7
0136	4	198.5	> 320	>1.6
2862	20	198.5	32	1.6
2717	20	195	320	1.6
	8	64	100	1.6
2167	9	6.4	100	1.6
2193	32	64	> 100	>1.6
3691	31	6	100	1.6
3355	35	61	100	1.6
3765		20	32	1.6
3542	31		32 32	1.6
4802	46	20		
4339	42	20	32	1.6
1976	1	67	> 100	>1.5
1996	3	66	100	1.5
2985	27	209	320	1.5
2583	19	22	32	1.5
3861	40	21	32	1.5
4373	31	21	32	1.5
1729	45	66	100	1.5
5555	53	208	320	1.5
3788	35	21	32	1.5
3509	33	2.1	3.2	1.5
5557	53	2.2	3.2	1.5
2487	19	223	320	1.4
3204	29	2.3	3.2	1.4
2178	8	73	> 100	> 1.4
3581	32	6.9	10	1.4
5562	53	70	100	1.4
3805	35	6.9	10	1.4
2563	32	23	32	1.4
3867	41	22	32	1.4
4739	44	70	100	1.4
4454	45	7.1	10	1.4
3860	40	23	32	1.4
4120	40	230	320	1.4
4177	39	3.6	5	1.4

Table 23 (Cont'd)

AVS No.	Shipment	<u>IC</u> ₅₀	MTC	TL
3851	40	7.3	10	1.4
4316	42	24	32	1.4
1972	1	79.3	100	1.3
1973	1	25	32	1.3
2615	19	74	100	1.3
2093	6	76	100	1.3
2212	10	7.6	10	1.3
2984	27	25	32	1.3
3695	32	248	320	1.3
2716	22	2.5	3.2	1.3
4233	42	238	320	1.3
4049	37	7.8	10	1.3
4241	39	7.8	10	1.3
4427	44	25	32	1.3
4384	43	7.9	10	1.3
4047	37	7.9	10	1.3
4890	46	25	32	1.3
4085	37	254	320	1.3
5491	53	24	32	1.3
3797	35	7.6	10	1.3
3593	30	0.25	0.32	1.3
3496	33	25	32	1.3
1999	3	2.7	3.2	1.2
3689	32	82	100	1.2
2188	9	84	100	1.2
2232	10	84	100	1.2
2517	20	2.7	3.2	1.2
2138	5	270	320	1.2
2285	11	269	320	1.2
2812	24	0.02	0.032	1.2
3106	28	27	32	1.2
2739	22	28	32	1.2
3205	29	2.8	3.2	1.2
1850	32	261	320	1.2
3680	32	82	100	1.2
1901	33	82	100	1.2
3677	32	2.7	3.2	1.2
3589	32	27	32	1.2
4261	39	8.1	10	1.2
4192	39	8.2	10	1.2
3789	35	8.2	10	1.2
4216	41	85	100	1.2
3858	40	0.85	- 1	1.2
4437	44	8.7	10	1.2
1738	45	87	100	1.2
0202	4	298	> 320	>1.1
0228	1	28.4	32	1.1

Table 23 (Cont'd)

AVS No.	Shipment	<u>IC</u> 50	MTC	II
0302	1	0.3	0.32	1.1
1089	5	0.9	1.0	1.1
2718	22	279	320	1.1
2652	21	287	320	1.1
0065	2	29	32	1.1
2429	17	9.1	10	1.1
3131	28	9.2	10	1.1
2971	27	30	32	1.1
2189	9	303	320	1.1
4281	42	2.8	3.2	1.1
4127	40	89	100	1.1
4859	48	9.1	10	1.1
4197	39	94	100	1.1
3902	41	9.5	10	1.1
1730	45	306	320	1.1
3488	33	2.8	3.2	1.1
3499	33	9.1	10	1.1
3351	31	30	32	1.1
3919	34	9.3	10	1.1
2403	16	30	32	1.1

^a Compounds identified by their AVS numbers are listed in descending order from the most active to the least active. Assays were done in Vero cells. An active compound is defined here as having a therapeutic index (TI) of greater than 1.0. The drug concentration which reduced the mean plaque number of 50% (50% inhibitory dose, IC_{50}) was calculated using a regression analysis program for semilog curve fitting and is expressed in $\mu g/ml$. The minimum cytotoxic drug concentration (MTC) is the lowest drug concentration at which toxicity was observed. This is also expressed in $\mu g/ml$. The therapeutic index (TI) of a compound was determined by dividing the MTC by the IC_{50} .

4.1.9.2 Confirmatory Assays:

Some of the compounds were sent to us in more than one separate drug shipment. These compounds were tested more than once. Data from the confirmatory assays are summarized in Table 24. If a compound showed a Therapeutic Index of ≥ 1 , then it was considered a candidate for confirmatory testing. Out of 10 confirmatory assays, 9 compounds were confirmed active during this contract period. If sufficient amounts of compound is available, the other compounds in Table 23 should be retested for confirmation.

Table 24

Confirmatory Assays for Compounds Found
Active Against Pichinde Virus

AVS No.	Shipment	IC ₅₀	MTC ug/ml	II
0646	2 2	0.1 0.4	100 > 10	>681 >27
0148	2 2	0.2 0.7	100 > 100	589 > 151
0140	4 4	0.5 0.2	> 100 0.32	>208 1.5
0001	4 4	2.5 3.7	> 100 32	>40 8.7
0230	1	9.8 71	320 320	33 4.5
2506	21 21	1.4 4.8	10 10	7.4 2.1
0094	1 1 1	30 56 44	>100 100 100	>3.3 1.8 2.3
2980	25 25	1.4 0.86	3.2 1.0	2.4 1.1
3788	35 35	21	32 3.2	1.5
4241	39 46	7.8 18	10 10	1.3 0.7

Compounds identified by their AVS number are listed in ascending order by AVS number.

IC₅₀ = The minimum drug concentration (μ g/ml) that inhibited the CPE by 50%, calculated by using a regression analysis program for semilog curve fitting.

MTC = Minimum cytotoxic drug concentration is the lowest drug concentration at which drug toxicity was observed.

TI = The Therapeutic Index of a compound was determined by dividing the MTC by the IC_{50} .

4.1.10 Vesicular Stomatitis Virus (VSV)

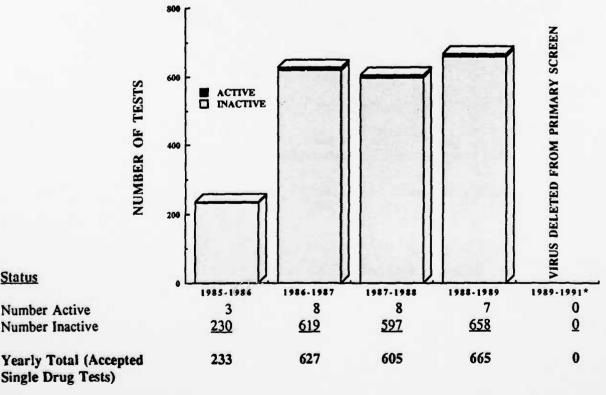
The number of single drug tests carried out against VSV during this contract period is summarized in yearly increments in Figure 43. During this five-year period two main in vitro antiviral assay protocols were implemented:

- A standard CPE inhibition assay by virus rating (VR) (Annual Report, December 15, 1988, Section 3.2.4).
- Since January, 1989, MTT based-antiviral assay format. 2.

A total of 2438 tests were performed during this contract period using both assay types. Routine testing was changed to the MTT-assay format to improve the efficiency and quality of the primary screening program in addition to being more cost-effective. Two positive control drugs (AVS-303 and 1160) were included in each assay. For the CPE-inhibition assay, the VR of AVS-303 ranged from 1.3 to 4.8, the IC₅₀ from 0.6 to 18.5 μ g/ml, and the TI from 1.1 to 17.9. The VR of AVS-1160 ranged from 1.6 to 4.9, the IC₅₀ from 0.2 to 2.3 μ g/ml, and the TI from 1.1 to 33.6. A wider range of values was obtained with the MTT assay. The TC₂₅ of AVS-303 ranged from 4.33 to 320, the IC₅₀ from 1.27 to 5.39 μ g/ml, and the SI from 1.94 to 73.07. The TC₂₅ of AVS-1160 ranged from 1.31 to 37.6, the IC₅₀ from 0.32 to 55.9 μ g/ml, and the SI from 1.07 to 36.03. Results of these positive controls (VR tests) were used as a guideline to assess the quality of each assay.

Out of the 2130 accepted single drug tests, 26 compounds demonstrated antiviral activity at greater than 50% reduction levels. This represents around 1.2% of the tested compounds having in vitro antiviral activity against VSV. The remainder, 2104 compounds (98.8%), were considered inactive with both assay protocols (Figure 43).

IN VITRO PRIMARY SCREEN: NUMBER OF COMPOUNDS FOUND ACTIVE AGAINST VESICULAR STOMATITIS VIRUS DURING THE CONTRACT PERIOD



Five-Year **Totals**

26

2104

2130

Status

Number Active

Number Inactive

Represents 14-month period (November 15, 1989 - January 31, 1991) Figure 43

As shown in Table 25 fourteen compounds demonstrated confirmed in vitro antiviral activity versus VSV during this contract period. The activity of the remaining compounds was either not confirmed on retest or there was not sufficient drug available for retest.

The three most active compounds identified during this contract period were AVS-5580 with a TAI of >73.5 and >59.5 and a SI of >1000 and >5060, AVS-5219 with a TAI of >71.99 and 28.27 and a SI of 127 and 10.6 and AVS-2350 with VR of 2.0 and a SI of 32 and 317. AVS-2503 also showed good antiviral activity with a VR of 4.1 and 4.8 and a SI of 46 and 75.

Table 25

Confirmatory Assays for Compounds Active Against Vesicular Stomatitis Virus

AVS	Ship-							Assay
No.	ment	Date	DIFRNTL	IC ₅₀	TC ₂₅	TAI	<u>SI</u>	Type
346ª	25	07/10/87	NA	0.41	1.00	0.95 ^b	2.42	CPE
	25	07/28/87	NA	0.09	0.32	1.40 ^b	3.60	CPE
1159	27	07/28/87	NA	15.95	100.00	1.90 ^b	6.27	CPE
	27	10/13/87	NA	8.86	100.00	2.60 ^b	11.29	CPE
1214	52	03/03/89	1.451	3.05	20.20	> 30.25	6.61	MTT
	52	05/12/89	1.477	2.33	5.78	10.52	2.48	MTT
1217	52	03/03/89	1.451	41.60	218.00	>29.06	5.24	MTT
	52	05/12/89	1.429	55.90	144.00	13.33	2.58	MTT
2350	14	11/06/87	NA	0.00	>20000.00	1.10 ^b	>0.00	CPE
	14	11/12/87	NA	632.00	200000.00	1.95 ^b	316.60	CPE
	14	02/04/88	NA	0.00	200000.00	0.00 ^b	0.00	CPE
	14	05/13/88	NA	6317.00	>200000.00	2.00 ^b	>31.70	CPE
2503	21	05/05/87	NA	2.17	100.00	4.10 ^b	46.10	CPE
	21	06/12/87	NA	1.33	100.00	4.75 ^b	75.36	CPE
2906	26	07/17/87	NA	7.80	< 100.00	2.10 ^b	< 12.81	CPE
	26	07/28/87	NA	8.86	100.00	2.60 ^b	11.29	CPE
3586	32	01/29/88	NA	1.80	32.00	3.70 ^b	17.70	CPE
	32	02/23/88	NA	1.60	32.00	4.30 ^b	20.10	CPE
3968	36	03/18/88	NA	89.50	320.00	1.25 ^b	3.58	CPE
	36	08/04/89	1.570	54.20	180.00	>17.98	3.32	MTT
4240	39	04/05/88	NA	8.00	32.00	2.60 ^b	4.00	CPE
	39	05/10/88	NA	11.80	< 100.00	1.70 ^b	< 8.50	CPE
4275	39	04/08/88	NA	0.22	1.00	3.60 ^b	4.49	CPE
	39	04/19/88	NA	0.16	0.32	4.10 ^b	1.95	CPE
	39	08/11/89	1.504	0.06	0.22	> 18.82	3.66	MTT
5219	51	03/03/89	1.315	1.96	249.00	>71.99	127.00	MTT
	51	05/12/89	1.477	1.85	19.60	28.27	10.60	MTT
5556	53	05/12/89	1.300	48.10	141.00	> 19.17	2.93	MTT
	53	06/02/89	1.391	41.80	155.00	24.99	3.70	MTT

Table 25 (Cont'd)

AVS No.	Ship- ment	Date	DIFRNTL	<u>IC</u> 50	TC ₂₅	TAI	SI	Assay Type
5580	54 54	05/05/89 05/19/89	1.153 1.657		> 100000. 00 > 100000. 00		> 1000.00 > 5060.00	MTT MTT

^a Compounds identified by their AVS numbers are listed in ascending numerical order. AVS-2350 and AVS-5580 were tested in Vero cells and activity is expressed as I.U./ml. All other assays were done in L929 cells.

DIFRNTL = The differential is the difference in the cell control and the virus control optical densities.

 IC_{50} = The minimum drug concentration ($\mu g/ml$) that inhibited CPE by 50%, calculated by using a regression analysis program for semilog curve fitting.

 TC_{25} = The minimum drug concentration (μ g/ml) that reduced cell viability by 25%.

TAI = Total Antiviral Index = the area between the cytotoxicity and the antiviral curves.

SI = Selective Index, calculated by dividing the TC_{25} by the IC_{50} .

NA = Not Applicable.

^b This value is a virus rating (VR) rather than TAI. The VR is a measurement of selective antiviral activity that takes into account the degree of inhibition of virus-induced CPE and the degree of cytotoxicity produced by the test compound similar to TAI.

4.1.11 Antiviral Screening In Vitro: Retroviruses:

The contract was modified in the second year (DAMD17-86-C-6013, modification No. P60001) to also include testing of the compounds supplied by USAMRIID against Human Immunodeficiency Virus (HIV) and other retroviruses. In April, 1987, the contract was amended to test 5,000 - 10,000 compounds per year against HIV for NCI. We used the XTT assay and automated plate readers. A computer was connected to the plate reader for data processing and then the data was down loaded to the NCI database by modem. This testing was later supported by two separate contracts (cooperative agreement No. DAMD17-88-H-8003 dated January 4, 1988 and NCI Contract N01-CM-87237 dated July 15, 1988). Results are presented below for antiviral testing done under this contract.

4.1.11.1 Primary Screening-HIV:

4.1.11.1.1 **USAMRIID Compounds:**

Approximately 500 AVS compounds supplied to us from USAMRIID were screened in the primary ATH8 cell assay for antiviral activity against HIV. Although the ATH8 human T-cell line proved to be very sensitive in detection of HIV antiviral activity, it was a troublesome cell line to maintain in quantity on a regular basis for use in assays.

Table 26 lists the most active AVS compounds, based on the therapeutic index and virus rating, identified in our primary screening. All of these compounds appeared to be very potent, with six of them, AVS-999, AVS-2353, AVS-2623, AVS-1764, AVS-206 and AVS-1603, having an ID₅₀ of 0.1 μ g/ml or lower. Several of these compounds appeared very active, having VR values greater than 4, with AVS-2576 having a VR of 5.2. Dideoxycytidine (ddC) was included with each set of assays as a positive control drug. It typically had a VR of 4 or greater and an ID₅₀ of <0.1 μ g/ml. A complete listing of all the USAMRIID compounds found to have some degree of activity in the primary HIV screen is presented in Appendix A, Table 9 of our second annual report dated December, 15, 1987.

Table 26
Compounds Active Against Human Immunodeficiency Virus^a

AVS#	Shipment	<u>VR</u>	VR*	ID ₅₀	MTC	II
0999	9	3.5	1.2	< 0.1	100	> 1000
2353	11	4.2	1.4	< 0.1	> 100	> 1000
2276	11	4.3	1.4	0.2	100	499
2623	19	3.1	1.0	< 0.1	32	>320
1764	22	3.1	1.0	< 0.1	32	> 320
2285	11	4.4	1.5	0.4	> 100	>254
1790 ·	21	2.6	0.9	0.5	100	184
2572	19	4.7	1.6	1.0	100	100
0206	9	2.0	0.7	0.1	10	100
1603	N/Ab	4.2	1.4	< 0.1	10	> 100
2576	19	5.2	1.7	1.0	100	100
2575	19	4.5	1.5	1.5	> 100	>69
2574	19	3.6	1.2	1.6	100	64
2567	21	3.4	1.1	1.8	> 100	> 54

a. Compounds identified by their AVS number are listed in descending order by activity. Assays were done in ATH8 cells.

b. AZT (not applicable).

- VR = A measurement of selective antiviral activity which takes into account the degree of inhibition of virus-induced CPE and the degree of cytotoxicity produced by the test compound, determined by a modification of the method of Ehrlich et al. (Ann. N.Y. Acad. Sci. 130:5, 1965).
- VR* = The designation for the virus rating calculated by the method of Sidwell and Huffman (Appl. Microbiol. 22:797, 1971).
- ID₅₀ = The minimum drug concentration (μ g/ml) that inhibited the CPE by 50%, calculated by using a regression analysis program for semilog curve fitting.
- MTC = Minimum cytotoxic drug concentration is the lowest drug concentration at which drug toxicity was observed.
- TI = The Therapeutic Index of a compound was determined by dividing the MTC by the ID_{50} .

Compounds exhibiting HIV antiviral activity in the initial ATH8 cell assay were retested in ATH8 cells to confirm the activity. The results of these confirmatory assays are presented in Table 27. Antiviral activity was confirmed for many of the compounds. Some of the best compounds, based on the VR and TI values, appeared to be AVS-999, AVS-2353, AVS-2358, AVS-2393, and AVS-2576. Compounds having confirmed activity were further evaluated in additional assays (see below).

Table 27

Confirmatory Assays for Compounds Active Against Human Immunodeficiency Virus^a

	Date						
AVS#	Shipment	Tested	<u>VR</u>	<u>VR*</u>	<u>ID</u> 50	MTC	TI
1	9	01/26/87	1.5	0.5	0.8	3.2	4.0
		07/08/87	2.0	0.7	-	10.0	-
206	9	01/29/87	2.0	0.7	0.1	10.0	100.0
		10/19/87	1.9	0.6	-	10.0	•
999	9	01/29/87	3.6	1.2	< 0.1	100.0	>1000.0
		03/16/87	1.9	0.6	2.2	32.0	15.0
		10/19/87	3.3	1.1	0.3	100.0	313.0
1790	21	07/16/87	2.6	0.9	0.5	100.0	184.0
		11/06/87	0.0	0.0		32.0	•
2274	12	03/11/87	1.2	0.4	0.3	3.2	10.0
		06/30/87	0.0	0.0	-	< 100.0	-
2275	12	03/16/87	1.0	0.3	-	10.0	-
		06/30/87	0.0	0.0	-	32.0	-
2276	11/	01/26/87	4.3	1.4	0.2	100.0	499.0
		06/16/87	0.0	0.0		<u><</u> 0.1	-
		06/25/87	0.5	0.2	-	10.0	-
		06/30/87	0.0	0.0	•	32.0	"II E
2278	11	02/13/87	1.4	0.5	1.8	100.0	56.0
		06/17/87	0.0	0.0	, , , ,	<u><</u> 0.1	-
2285	11	02/03/87	4.4	1.5	0.4	> 100.0	>253.0
		07/01/87	1.7	0.6		100.0	
2332	12	03/11/87	1.7	0.6		> 100.0	
		06/25/87	1.8	0.6		> 100.0	
		07/01/87	1.1	0.4	-	>100.0	
		07/08/87	2.0	0.7	5.7	100.0	18.0
2340	12	03/19/87	1.5	0.5	9.5	> 100.0	>11.0
		07/01/87	1.3	0.4	•	> 100.0	+

Table 27 (Cont'd)

AVS#	Date Shipment	Tested	<u>VR</u>	VR*	<u>ID</u> 50	MTC	п
2353	11	01/26/87	4.2	1.4	< 0.1	> 100.0	> 1000.0
	-	06/16/87	0.0	0.0	-	< 0.1	-
		07/01/87	2.2	0.7	12.1	>100.0	> 8.2
2358	15	03/18/87	1.1	0.4	< 0.1	1.0	10.0
		07/01/87	1.2	0.4	43.0	> 100.0	>2.4
2365	15	03/22/87	1.8	0.6	L	> 100.0	-
		06/30/87	1.5	0.5	23.0	100.0	4.4
		07/01/87	0.5	0.2	-	>100.0	-
		07/08/87	1.3	0.4		100.0	-
2393	16	07/22/87	1.9	0.6	0.6	32.0	57.0
		10/19/87	3.3	1.1	59.0	100.0	1.7
2397	16	07/22/87	1.1	0.4	17.0	32.0	1.9
		10/19/87	2.4	0.8	-	32.0	-
2404	16	05/15/87	1.0	0.3	- [1]	10.0	
		07/24/87	0.0	0.0	-	10.0	-
2407	16	05/15/87	1.7	0.6	0.9	10.0	11.0
		07/06/87	0.0	0.0	-	32.0	-
2408	16	05/15/87	1.1	0.4	-	< 100.0	
		07/06/87	0.0	0.0	-	100.0	•
2488	19	06/01/87	2.4	0.8	3.2	32.0	10.0
		07/09/87	0.3	0.1	-	32.0	-
2572	19	06/05/87	4.8	1.6	1.0	100.0	100.0
		07/09/87	0.0	0.0	-	10.0	-
		10/29/87	0.0	0.0	-	100.0	
2574	19	06/05/87	3.6	1.2	1.6	< 100.0	64.0
		07/09/87	0.6	0.2	-	10.0	- 1 -
2576	19	06/05/87	5.2	1.7	1.0	100.0	100.0
		06/17/87	3.1	1.0	2.4	100.0	41.0
2581	19	06/05/87	3.4	1.1	3.2	100.0	31.0
		07/10/87	0.4	0.1	-	10.0	•
2621	19	07/08/87	1.7	0.6	-	32.0	
	• •	07/23/87	2.4	0.8	11.2	100.0	8.9

Table 27 (Cont'd)

43704	Date	m	170	77D.#	TD.) /TP/C	mr.
AVS#	Shipment	Tested	<u>VR</u>	VR*	ID_{50}	MTC	II
2708	22	10/14/87	1.3	0.4	0.1	1.0	10.0
		10/19/87	2.5	0.8	-	32.0	•
2748	22	05/20/87	1.1	0.4	0.1	1.0	8.2
		07/10/87	0.0	0.0	•	<u><</u> 0.1	-
2749	22	05/20/87	1.9	0.6	1.8	32.0	18.0
		06/17/87	0.0	0.0	-	10.0	-
		07/22/87	0.0	0.0	-	100.0	-

- a. Compounds are listed in ascending numerical order by AVS number. The results from the primary and the subsequent confirmatory CPE-inhibition assays are listed chronologically for each compound.
- VR = A measurement of selective antiviral activity which takes into account the degree of inhibition of virus-induced CPE and the degree of cytotoxicity produced by the test compound, determined by a modification of the method of Ehrlich et al. (Ann. N.Y. Acad. Sci. 130:5, 1965).
- VR* = The designation for the virus rating calculated by the method of Sidwell and Huffman (Appl. Microbiol. 22:797, 1971).
- ID₅₀ = The minimum drug concentration (μ g/ml) that inhibited the CPE by 50%, calculated by using a regression analysis program for semilog curve fitting.
- MTC = Minimum cytotoxic drug concentration is the lowest drug concentration at which drug toxicity was observed.
- TI = The Therapeutic Index of a compound was determined by dividing the MTC by the ID_{50} .

Further assays of AVS-999 were conducted on the following samples of the drug: (1) purified samples of 4-acetyl-4-phenyl-piperidine (PAP-base and PAP-HC1), purified from inactive drug received from the NCI (NSC 613291); (2) original and new samples obtained from Technassociates, and (3) a drug sample obtained from Dr. Raymond Schinazi (VA Medical Center, Decatur, GA). The samples were assayed in parallel on ATH8 cells and scored by visual assessment of CPE. The results, summarized in Table 28, showed that all of these samples were active, which was reflected in their VR scores of 2.5 - 3.3, although the original sample and the PAP-HC1 appeared to be less toxic, resulting in higher TI values for those two samples. It should be noted from the data that the maximum reduction in CPE observed at any concentration level in this set of assays was 58%, in contrast to parallel data for ddC in which 100% reduction was observed supporting the lack of concentration-dependency and the marginal nature of the results for AVS-999. The new samples obtained from Technassociates was identical by HPLC profile to the material received from the NCI (NSC 613291) and it contained the same impurities. This material was significantly less active than the purified compound and the original sample submitted as AVS-999.

Table 28

Retesting of AVS-999 Samples^a

AVS			ID_{50}	MTC	Therapeutic
No.	Sample	<u>VR</u>	$(\mu g/ml)$	(µg/ml)	Index
999	PAP-Base	3.2	3.2	100	31.25
999	PAP-HC1	2.7	0.32	> 100	>312.50
999	Schinazi Sample	3.1	3.2	100	31.25
999	Original Sample				
	(Technassociates)	3.3	0.32	100	312.50
999	New Sample				
	(Technassociates)	2.5	>32.0	100	< 3.13
2639	ddC (Positive Control				
	drug)	3.8	0.1	10	100

⁼ Assays were done in ATH8 cells.

VR = A measurement of selective antiviral activity which takes into account the degree of inhibition of virus-induced CPE and the degree of cytotoxicity produced by the test compound, determined by a modification of the method of Ehrlich et al. (Ann. N.Y. Acad. Sci. 130:5, 1965).

VR* = The designation for the virus rating calculated by the method of Sidwell and Huffman (Appl. Microbiol. 22:797, 1971).

ID₅₀ = The minimum drug concentration (μ g/ml) that inhibited the CPE by 50%, calculated by using a regression analysis program for semilog curve fitting.

MTC = Minimum cytotoxic drug concentration is the lowest drug concentration at which drug toxicity was observed.

TI = The Therapeutic Index of a compound was determined by dividing the MTC by the ID_{50} .

4.1.11.1.2 NCI Compounds:

During this period all elements of the XTT (tetrazolium dye) assay system were brought into operation. During the short period that this Task was included in this contract, we tested approximately 1700 NSC compounds supplied by NCI against HIV in MT-2 cells of this number, 20 compounds were considered active or possibly active and the remainder were either inactive or represented unsatisfactory tests. The most frequent cause of unsatisfactory tests was poor infectivity of virus. One drug, NSC-614846, demonstrated significant activity comparable to dideoxycytidine (ddC, NSC-606170, AVS-2639). The results for ddC are shown in Figure 44 and results for NSC-614846 assayed in MT-2 cells, are shown in Figure 45. Inoculation was by cocultivation with H9/IIIB cells (20 cells/well). This compound (NSC 614846) had an ID₅₀ of 0.009 μ g/ml, a 50% cytotoxicity concentration (TC₅₀) of 3.8 μ g/ml, and a TI (calculated as the TC₅₀/ID₅₀) of 404. Figure 46 is an example of the data sheet resulting from this assay.

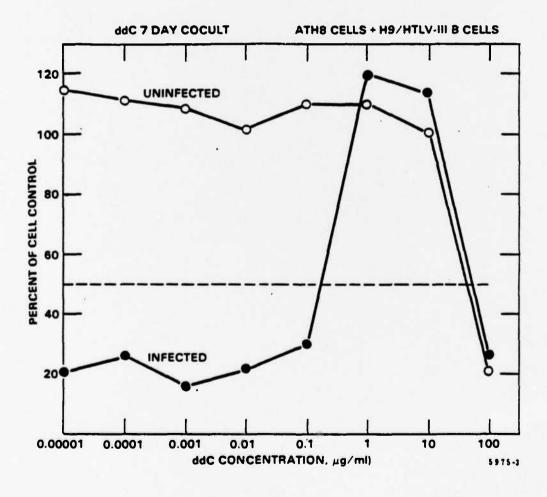
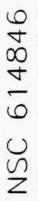
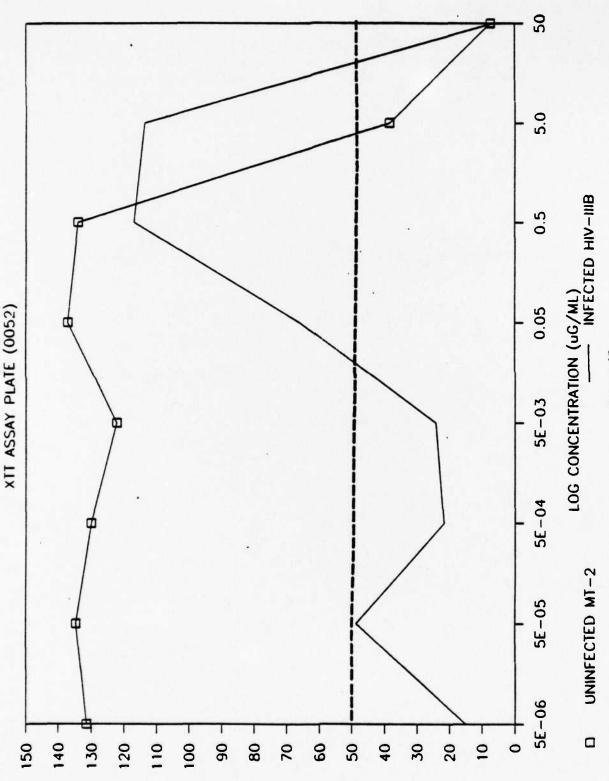


Figure 44

Results from an XTT assay of the antiviral activity of 2',3'-dideoxycytidine against HIV in ATH8 cells.





S OF CONTROL

Figure 45
Results from an XTT assay of the antiviral activity of NSC-614846 against HIV in MT-2 cells.

EXPERIMENT ID : 9N 0052 REPORT DATE : 09-FEB-88 PLATE BARCODE CONFIG: 017 ASSAY : 0052 : XIT INCC DATE : 03-DEC-87 DRUG ADD DATE : 03-DEC-87 : 03-DEC-87 MR600 READ DATE : 03-DEC-87 STAIN DATE OPTICAL DENSITIES 01 02 03 04 05 06 07 08 10 11 12 0.017 0.018 0.017 0.221 0.216 0.000 0.220 0.219 0.222 0.291 0.236 0.020 0.018 0.232 0.692 0.396 0.588 0.455 0.595 0.986 1.429 1.525 0.297 0.018 B 0.018 0.231 0.667 0.393 0.949 0.482 0.398 0.944 1.639 1.471 0.304 0.019 C 0.020 0.217 0.419 0.793 0.284 0.309 0.627 1.298 1.299 1.484 0.532 0.019 D 0.020 0.223 1.328 0.756 0.407 0.719 0.410 1.510 1.564 1.558 1.000 0.018 E 0.019 0.223 1.353 1.696 1.732 1.680 1.593 1.759 1:726 0.656 0.308 0.018 0.019 0.225 1.351 1.350 1.224 1.276 1.403 1.377 1.503 1.623 1.281 0.017 G 0.020 0.020 0.020 0.220 0.210 0.000 0.219 0.219 0.211 0.214 0.210 0.017 CALCULATED RESULTS (INFECTED) WELL GRP NSC TTYPE CONC ∞ MEAN SD CV T/C% IND EC50/90/100 CELL LINE : MT-2 SAMPLE: 0100 E03-G03 VEH 54 1.119 0.014 0.010 1 BACKGD 0.225 0.006 0.025 B02-G02 B04-C04 614846 11 5.00E-06 u 1 0.169 0.002 0.005 15.1 N 9.38E-03 B05-C05 0.543 0.255 0.332 48.6 614846 5.00E-05 u 66 N 1.47E-01 11 5.00E-04 u 1 0.041 21.7 B06-C06 0.243 0.019 * 2.31E-01 614846 11 5.00E-03 u 66 B07-C07 614846 11 0.271 0.139 0.281 B08-C08 614846 5.00E-02 u 1 0.740 0.030 0.031 66.1* 11 B09-C09 11 614846 5.00E-01 u 1 1.309 0.148 0.097 117.0 B10-C10 614846 0.038 11 5.00E+00 u 1 1.273 0.025 113.8 B11-C11 0.005 11 5.00E+01 u 1 0.075 0.016 614846 6.7 B03-D03 HIVENIL 66 0.368 0.151 0.151 32.8 CALCULATED RESULTS (UNINFECTED) WELL GRP NSC TTYPE CONC MEAN SD CV T/C% IND IC50/90/100 ∞ CELL LINE : MT-2 SAMPLE: 0100 E03-G03 1 1.119 0.014 0.010 **VEH 54** F04-F04 lU 5.00E-06 u 1 1.471 0.000 0.000 131.5 614846 3.79E+00 F05-F05 614846 1U 5.00E-05 u 1 1.507 0.000 0.000 134.7 4.12E+01 F06-F06 614846 5.00E-04 u 1 1.455 0.000 0.000 130.0 >5.00E+01 1U F07-F07 5.00E-03 u 1 0.000 0.000 122.3 614846 lU 1.368 F08-F08 614846 1U 5.00E-02 u 1 1.534 0.000 0.000 137.1 F09-F09 614846 5.00E-01 u 1 1.501 0.000 0.000 134.1* lU F10-F10 614846 lU 5.00E+00 u 1 0.431 0.000 0.000 38.5* F11-F11 614846 0.000 lU 5.00E+01 u 1 0.083 0.000 In Vitro Therapeutic Indices IC50/EC50 IC90/EC90 IC100/EC100 614846 * 4.04E+02 > 2.79E+02 > 2.16E+02

Figure 46
Example data sheet form an XTT assay.

4.1.11.3 Secondary Evaluations (FAIDS, MAIDS, and SAIDS):

4.1.11.3.1 Feline Leukemia Virus - FAIDS Variant:

Initially, several compounds were screened in parallel using both the FeLV (FAIDS) assay and the HIV ATH8 assay, and occasionally, such as when the ATH8 assay was not working properly, the FeLV (FAIDS) assay was used as the primary screen for retrovirus antiviral activity. Four additional compounds were identified having antiviral activity in this focus-inhibition assay (Table 29). The most active compound assayed in this model was AVS-2708. It had a VR of 4.6, and ID₅₀ of 1.2 μ g/ml, and a TI of 82.5. This compound also showed antiviral activity against HIV (VR values of 1.3 and 2.5, Table 27). Compounds AVS-2280, AVS-2291, and AVS-222 did not show activity against HIV.

Table 29

Compounds Active Against Feline Leukemia Virus FAIDS Variant*

AVS#	Shipment	<u>VR</u>	VR*	ID ₅₀	MTC	I
2708	22	4.6	1.5	1.2	100	82.5
2280	11	3.2	1.1	4.5	100	22.1
2291	11	1.7	0.6	4.6	32	7.0
0222	9	1.4	0.5	31.4	32	1.0

- a. Compounds identified by their AVS number are listed in descending order by activity. CPE-inhibition assays were done in F81C cells.
- VR = A measurement of selective antiviral activity which takes into account the degree of inhibition of virus-induced CPE and the degree of cytotoxicity produced by the test compound, determined by a modification of the method of Ehrlich et al. (Ann. N.Y. Acad. Sci. 130:5, 1965).
- VR* = The designation for the virus rating calculated by the method of Sidwell and Huffman (Appl. Microbiol. 22:797, 1971).
- ID₅₀ = The minimum drug concentration (μ g/ml) that inhibited the CPE by 50%, calculated by using a regression analysis program for semilog curve fitting.
- MTC = Minimum cytotoxic drug concentration is the lowest drug concentration at which drug toxicity was observed.
- TI = The Therapeutic Index of a compound was determined by dividing the MTC by the ID₅₀.

Five compounds have been assayed in the FeLV (FAIDS) focus-inhibition assay for confirmation of antiviral activity. These results are summarized in Table 30. Four of the compounds, AVS-0001, AVS-999, AVS-2285, and AVS-2353, showed antiviral activity in this *in vitro* feline retrovirus model; AVS-2358 did not display activity in this assay. AVS-2639 (dideoxycytidine) was the positive control compound for this assay and yielded VR's in the range 1.0 - 3.4, ID₅₀'s in the range $0.3 - 3.6 \mu g/ml$, and TI values of 2.8 - 32.4.

4.1.11.3.2 Simian retrovirus - SAIDS:

Three compounds were assayed in the SAIDS syncytia-inhibition assay during the reporting period. Because of the nature of this assay, a virus rating was not calculated for test compounds. Thus, antiviral activity of a compound was determined by a therapeutic index of 1.0 or greater. Two compounds, AVS-1 and AVS-2285, showed antiviral activity according to this scheme. AVS-999 was inactive in this assay. The results are summarized in Table 30.

4.1.11.3.3 Murine AIDS (MAIDS:

Two drugs which demonstrated significant antiviral activity in the HIV primary screen AVS-2639 (ddC) and AVS-01 (Ribavirin), have been tested in the CAS-BR-M murine leukemia virus plaque-reduction assay. AVS-2639 (ddC) was the more active of the two drugs tested. Only slight toxicity was observed at 100 μ g/ml (Figure 47-B) and 100% plaque-reduction was observed at drug concentrations as low as 10 μ g/ml (Figure 47-A). The MIC₅₀ was 1.8 μ g/ml and the TI was 54.5. This drug will be used as the positive control drug in future drug assays.

AVS-01 (Ribavirin) also demonstrated antiviral activity in this assay (Figure 48-A). The drug was toxic to the SC-1 cells at 32 μ g/ml and partially toxic at 10 and 3.2 μ g/ml (Figure 48-B). The MIC₅₀ was 0.5 μ g/ml and the TI was 5.9. AVS-2639 was included in the assay as a positive control drug. It had an MIC₅₀ of 1.2 μ g/ml and a TI > 8.2.

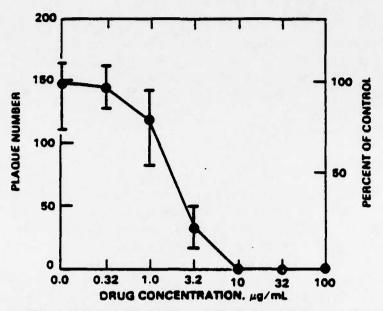


Figure A. Effect of AVS 2639 (ddC) on CAS-Br-M replication in SC-1 cells.

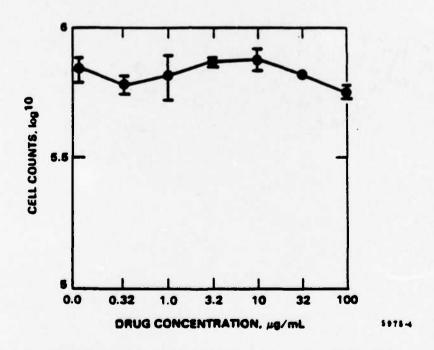


Figure B. Effect of AVS 2639 (ddC) on host cell multiplication.

Figure 47

Antiviral activity of 2',3'-dideoxycytidine in the in vitro murine retrovirus assay.

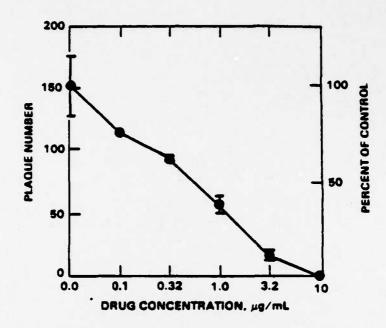


Figure A. Effect of AVS 1 (ribavirin) on CAS-Br-M replication in SC-1 cells.

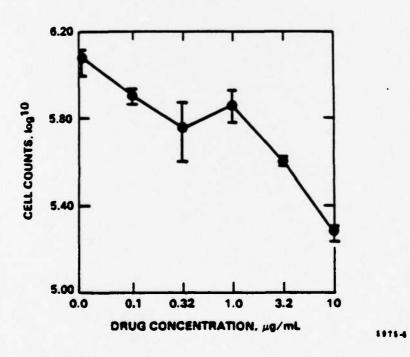


Figure B. Effect of AVS 1 (ribavirin) on host cell multiplication.

Figure 48

Antiviral activity of Ribavirin in the in vitro murine retrovirus assay.

Table 30

Secondary Testing of AVS Compounds Showing Confirmed Activity Against HIV in ATH8 Cell Assays*

1.0 26/320 12 0.55/3.2 5.9 0.2/32 139 K 6.8 -/>320	26/320 12 0.55/3.2 5.9 0.2/32 139 X -/>320		PAIDS ID50/NTC	2	SAIDS IDSO/NTC TI	NAIDS IDSO/NTC TI	HeLa-T4 ⁺ Syncytia Assay ID50/NTC TI	Fluorescence IDSO/MTC TI
-/>320	-/>320	3.2/3.2	-	0				×
28/>320 >11	28/>320 >11	14.8/100 6.8 -/ 32 - 11.5/>32 >2.8 3.9/>10 >2.6	6 % %			×	^	
X -/>100 X X X 2.7/32 12 X X X X X X X X X X X X X X X X X X X X X	X	29.8/ 32 1	-	-:	28/>320 >11	×		×
X	X X 2.7/32 12 X X X X X X X X X X X X X X X X X X X	×			×	×		×
x x x x x x x x x x x x x x x x x x x	x x x x x x x x x x x x x x x x x x x	×			×	×		×
* * * *	x x x x x x x x x x x x x x x x x x x	9.7/100		10	×	×		
* * *	x x x x x x x x x x x x x x x x x x x	-/10	·	1	×	×	×	×
* *	x x x x x x x x x x x x x x x x x x x	×			×	×	×	×
×	4.1/>10 >2.4 1.2/>10 >8.2 <0.1/32 >320	×			×	×	×	×
	4.1/>10 >2.4 1.2/>10 >8.2 <0.1/32 >320	×			×	×	, *	*
		0.68/10		15	4.1/>10 >2.4	1.2/>10 >8.2		0.48/>3.2 >8.7

a. Results are shown for compounds evaluated in five different secondary assays. The abbreviations are defined in the footnote to Table 25. Dideoxycytidine (ddC, AVS-2639) was assayed in parallel as a positive control drug. The "X" indicates assays scheduled for testing.

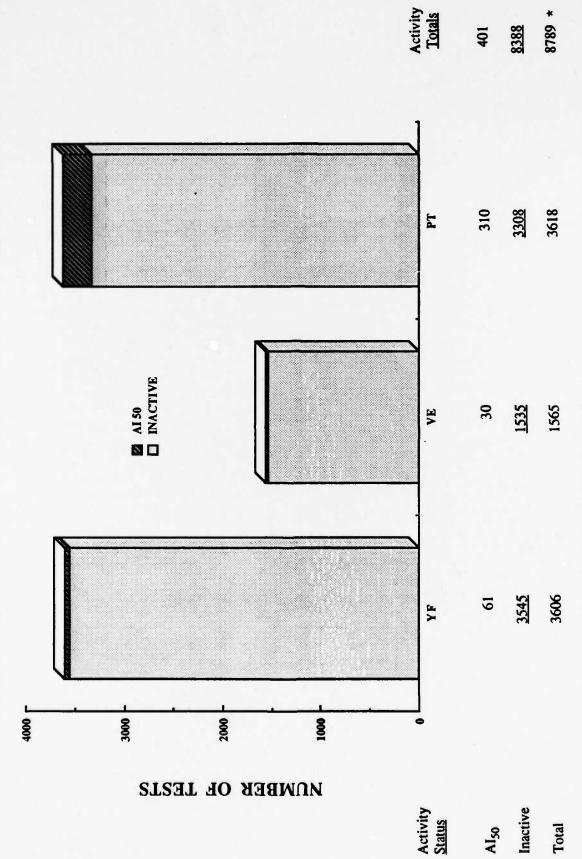
4.2 In Vitro Prescreen Antiviral Evaluation

Large numbers of plant extracts were available for screening for *in vitro* antiviral activity. We were requested to develop an assay that would: 1) allow more compounds to be evaluated per microtiter plate than in the regular antiviral screen and 2) pre-evaluate plant extracts as well as other extracts of natural origin for activity against three indicator virus families. The three viruses selected by USAMRIID for this purpose were the attenuated, vaccine strain (17D) of Yellow Fever virus (YF), Punta Toro virus (PT), and Venezuelan Equine Encephalomyelitis virus (VE). The prescreen should select candidate compounds for screening against the Asibi strain of YF and the other more virulent target viruses in the full screen.

A prescreen procedure was developed which utilized MTT and evaluated five compounds per virus per 96-well plate. Vero cells were seeded as monolayer cultures in COSTAR 96-well plates at 18,000/0.2 ml/well in MEM + 10% heat-inactivated fetal bovine serum (Δfbs). The plates were incubated approximately 24 hours prior to use. From the time that prescreen assays were started on this contract (June 1989 through January 31, 1991), a total of 3809 test compounds were received for evaluation in the prescreen antiviral assay. Approximately 14,000 in vitro antiviral prescreen assays (tests) were performed during this contract period (5667 tests with YF, 2235 tests with VE and 6026 tests with PT). It is worthwhile to note that we added the VE virus to the prescreen in June, 1990 therefore fewer assays were tested with this virus.

Out of 8789 accepted single drug tests from all three viruses, 401 demonstrated antiviral activity at \geq 50% reduction levels. This represents \sim 5% of the tested natural products having in vitro antiviral activity against these viruses (YF, VE, and PT). The remainder 8388 (95%) are to be considered inactive with the prescreen assay protocol. The antiviral activity results of the prescreen assays are summarized in Figure 49. In reality we have confirmed some compounds with \leq 20% original antiviral activity, i.e., if the cytotoxicity was zero or very minimal at 1000 μ g/ml. Based upon the present prescreen confirmatory criteria, the correlation between prescreen actives and confirmed actives from the primary screen was 67% (Table 34). The detailed results are summarized in the following sections for each virus.

ACTIVE COMPOUNDS FROM THE PRESCREEN ASSAY



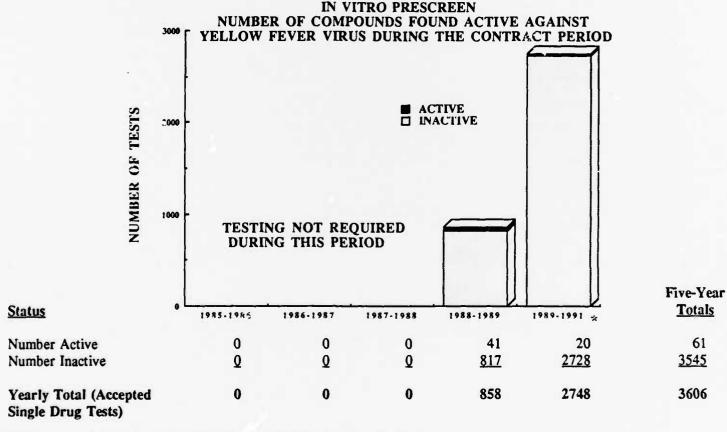
* Grand Total of Accepted Single Drug Tests (Excluding Positive Control Tests)

Figure 49

4.2.1 Prescreen Assay (Yellow Fever Virus [YF]):

The number of single drug tests carried against YF during this contract period is summarized in yearly increments in Figure 50. During this nineteen-month period (June, 1989 - January, 1991) 5667 tests were performed against YF-virus with MTT-assay format. Out of these, 266 were control compound assays-selenazofurin (AVS-0253) being the primary positive control compound. Nine hundred thirty-eight (938) tests were internal (+ + +) virus load, cell load, and other quality control tests. Eight hundred fifty-seven (857) tests were considered unsatisfactory based on the preliminary criteria of the quality controls set during this reporting period. The rest, totaling 3606 were actual single drug tests. The 857 unsatisfactory tests represent a 15% rejection rate based on present preliminary quality control parameters used for the YF-virus.

Out of the 3606 accepted single drug tests, 61 compounds demonstrated antiviral activity at greater than 50% reduction levels. This represents a 1.7% of the tested compounds having in vitro antiviral activity against YF-virus. The remainder, 3545 compounds (98.3%), are to be considered inactive with both assay protocols (Figure 50).



Represents 14-month period (November 15, 1989 - January 31, 1991)
Figure 50

- 4.2.1.1 <u>Prescreen YF-Quality Controls:</u> Two positive control compounds (Selenazofurin and 2-Thio-6 Azauridine) were used in the daily assay sets as antiviral activity quality controls. The antiviral performance of the unknown compounds is compared to that of the positive control compounds. Compounds with equal to or better antiviral potency are considered active and are worthy of further in vitro profile studies and in vivo testing.
- 4.2.1.1.1 <u>Antiviral Activity of Selenazofurin vs YF Virus:</u> A summary of the antiviral and cytotoxicity performance of the primary control compound, AVS-0253 (Selenazofurin) is presented in Figure 51-A for 120 tests performed during June, 1989 through January, 1991.

<u>Control Compound-Antiviral Performance:</u> Selenazofurin (AVS-0253) has been the sole control compound against YF in these MTT-assay prescreens. The mean and median antiviral inhibition and cytotoxicity patterns of the positive control drug (Selenazofurin) are illustrated in Figure 51-A.

The 120 control tests performed with Selenazofurin gave a mean Selectivity Index (SI) of 1.60 (SD \pm 3.10) and the median value was 0, indicating poor antiviral selectivity for Selenazofurin. The reason for this discrepancy is that Selenazofurin does not consistently reach the 50% antiviral reduction level, thus SI calculations cannot be executed properly. (SI is calculated by dividing the TC_{25} by the IC_{50}).

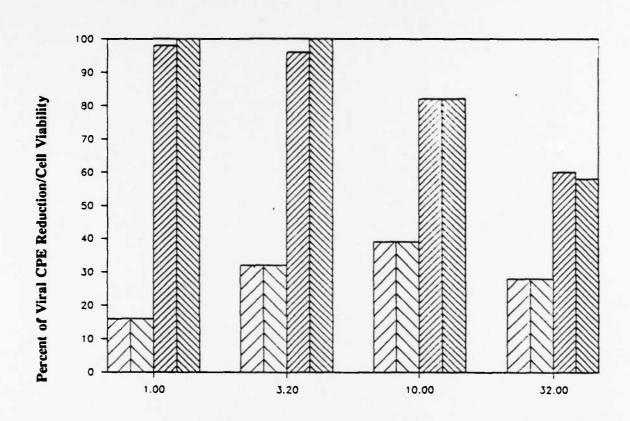
The mean Antiviral Inhibitory Concentration 50% (IC₅₀) was 1.53 μ g/ml (SD \pm 2.70). The median IC₅₀ value was 0 μ g/ml. This indicates that Selenazofurin does not consistently reach 50% antiviral reduction levels, at any tested drug concentration level. During this reporting period, the highest concentration of Selenazofurin was varied from 1 to 320 μ g/ml to properly evaluate the maximum antiviral effect and cytotoxicity pattern of Selenazofurin.

The maximum antiviral inhibitory level of 120 Selenazofurin tests (Figure 51-A) varies depending upon which drug concentration scale is used. Maximum antiviral effect ($\sim 39\%$) was found with a simultaneous $\sim 20\%$ cytotoxic suppression. Above 10 μ g/ml concentration Selenazofurin starts to lose its antiviral potency with increasing cytotoxicity. An increase of the concentration of Selenazofurin to 320 μ g/ml does not improve the antiviral activity (Figure 51-A). Actually antiviral activity decreases from 32% (at 3.2 μ g/ml) to $\sim 28\%$ (at 32 μ g/ml).

As previously reported in the seventh semiannual report, a different maximum antiviral value is obtained depending upon which Selenazofurin concentration scale is used. This scale measures more accurately the maximum antiviral effect of the control compound selenazofurin ($10 \mu g/ml$). In those tests, the \log_{10} scale of 0.1 - $100 \mu g/ml$ measured more accurately the antiviral effect of Selenazofurin, whereas the \log_{10} scale of 0.32 - $320 \mu g/ml$ more accurately measured the cytotoxicity effects.

In the present, 120 control assays we tested Selenazofurin at 0.5 \log_{10} scale concentrations (1 - 32 μ g/ml) to maximize the correct measurements of its antiviral and cytotoxicity effects. This enabled us to monitor our quality control parameters more accurately.

SELENAZOFURIN -VS- YF VIRUS (PRESCREEN PROTOCOL)



CONCENTRATION (µg/ml)

Mean %Viral CPEReduction			Median % Viral CPE Reduction		ZZZ Mean Cell Vial		Media Cell Vi	
	% Vir	al CPE Re	duction			% Cell	Viability	
Conc. (µg/ml)	1	3.2	10	32	1	3.2	10	32
Mean	16	32	39	28	98	96	82	60
Median	16	32	39	28	100	100	82	58
Std. Dev.	0.10	0.15	0.18	0.17	0.03	0.07	0.14	0.17

Figure 51-A
Average Antiviral and Cytotoxicity Values for 120 Positive Control Compound Assays

4.2.1.1.2 <u>Maximum Antiviral Effect of Selenazofurin vs YF Virus:</u> Since the metabolic activity of the cells was an unknown function during the testing period, it was monitored indirectly by measuring the maximum antiviral effect of the control compound Selenazofurin. This demonstrated the amount of infectious virus that was produced by the cells (Maximum Percent CPE).

A bar graph scatter plot (Figure 52-A) depicts the distribution of the maximum antiviral reduction values 120 control compound prescreen assays for Selenazofurin. The results indicate that the average maximum antiviral reduction obtained with the present SOP is around 45% (SD \pm 14.60) reduction levels. The maximum reduction levels vary from 25 - 93% but remain quite consistently around the median of 43%. The assay control values give a shifted bell-shaped distribution curve toward the median 32% reduction level. This indicates quite a consistent day-to-day performance of the control compound in the YF prescreen-MTT assay.

During this period the positive control compound performance criteria for Selenazofurin versus the YF virus in the pre-screen format has not been set to a definite endpoint. We have collected data in order to find out what would constitute a reasonable quality control endpoint and the data indicates that it could be set at the 25% reduction level. In order to measure the maximum antiviral endpoints of Selenazofurin correctly, the concentration scale and the highest concentration to be used, must be evaluated at the proper (semi-log) scale as seen in Figure 51-A.

Selenazofurin is active in vitro against YF virus and functions as a reasonable quality control compound. On the other hand, regardless of the performance of the YF-quality control drug Selenazofurin, around 20 other compounds have equal to or better antiviral activity against YF virus than AVS-0253.

Variation of the Maximum Antiviral Effect YF Virus - VS - Selenazofurin (Prescreen Protocol)

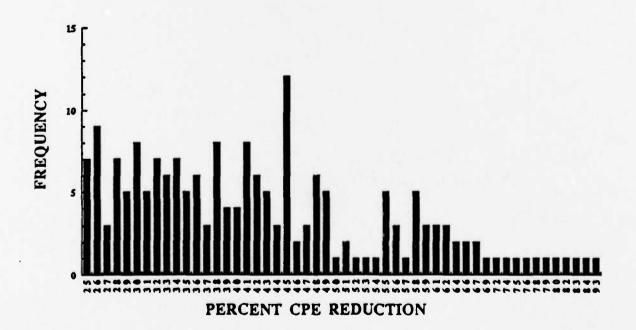


Figure 52-A

Maximum Antiviral CPE Reduction (%).

Summary of 120 Control Tests.

4.2.1.1.3 Cellular Cytotoxicity of Selenazofurin vs YF Virus:

<u>YF-Control Compound-Cytotoxicity Performance:</u> The 120 cytotoxicity values of the positive control compound Selenazofurin are also very consistent. The mean cell Toxic Concentration 25% (TC₂₅) was 17.90 μ g/ml (SD \pm 9.00) and the median was 16.05 μ g/ml (range of 2.81 - 32 μ g/ml).

As can be seen in Figure 51-A, a definite TC_{25} toxicity value can be measured with 32 μ g/ml 0.5 log_{10} scale. Further increase in the concentration of Selenazofurin would be needed to consistently evaluate the maximum cytotoxicity of Selenazofurin.

Also, Figure 51-A indicates that when the cytotoxicity reaches $\sim 20\%$ at 10 μ g/ml, the control compound (Selenazofurin) has reached simultaneously its maximum antiviral effect. The cytotoxic effect of Selenazofurin is insignificant below 3.2 μ g/ml. The average cytotoxicity reached $\sim 40\%$ at 32 μ g/ml, which was the highest Selenazofurin concentration in most tests.

Selenazofurin has a definite cytotoxic suppression on cellular metabolism and growth. The TC_{25} toxicity can be achieved with the 32 μ g/ml concentration of Selenazofurin. Therefore, a readjustment to 100 μ g/ml (as being the highest Selenazofurin concentration tested) is not needed. However, at this concentration (32 μ g/ml) the TC_{50} cannot yet be measured consistently.

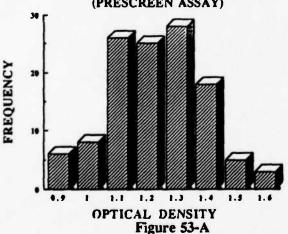
4.2.1.1.4 YF-Assay Plate Quality Controls: Cell Load and Virus Load Parameters (Selenazofurin): The MTT assay is fundamentally dependent upon the quality of the assay plates. Our large-scale antiviral testing is dependent upon the uniformity of the test plates produced for the daily assays. Equal numbers of cell load and virus load as well as the consistent performance of the reagents used daily was monitored. A sample of the plate variation control for the period of November, 1989 through January, 1991, is presented in Figures 53-A, 54-A and 55-A.

YF-Control Compound-Cell Load Performance: A bar graph scatter plot of the mean cell control (O.D. reading) of 120 control assays is plotted in Figure 53-A. The results indicate that the cell O.D. readings reached a mean 1.228 (SD \pm 1.55) with a median of 1.233 (range of 0.853 - 1.570). This indicates that a uniform and equal number (18,000 cells/well) of cells are being loaded into every well in the 96-well plate during the day-to-day operation. The cells reduced MTT to formazan giving maximum blue color uniformly and consistently.

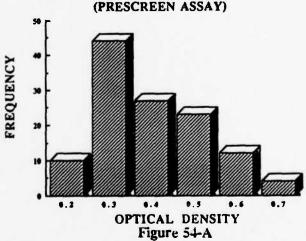
<u>YF-Control Compound-Virus Load Performance</u>: A bar graph scatter plot of the mean virus load O.D. readings of the 120 control assays is presented in Figure 54-A. The results indicate that the average virus load O.D. reading is 0.397 (SD \pm 0.122) with a median of 0.379 (0.155 - 0.697). This demonstrates that a reasonable cell destruction is taking place and a uniform load of virus (32 TCID₅₀) is administered on the cell monolayer with consistent viral CPE results.

YF-Control Compound-Assay Differential Performance: A bar graph scatter plot of the mean O.D. differential values of the 120 control assays is provided in Figure 55-A. The results indicate that the average differential O.D. reading is 0.831 (SD \pm 0.152) with a median of 0.821 (range 0.516 - 1.256). The single bell-shaped curve is reasonably sharp and uniform. This reflects that the assays are executed consistently and are repeatable during day-to-day operation with close to 83% measurement accuracy.

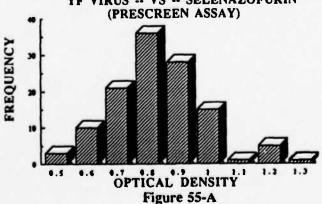
VARIATION OF THE CELL (LOAD) CONTROLS
YF VIRUS -- VS -- SELENAZOFURIN
(PRESCREEN ASSAY)



VARIATION OF THE VIRUS (LOAD) CONTROLS YF VIRUS -- VS -- SELENAZOFURIN (PRESCREEN ASSAY)



VARIATION OF THE TEST DIFFERENTIAL YF VIRUS -- VS -- SELENAZOFURIN



4.2.1.1 Prescreen YF-Ouality Controls:

4.2.1.1.1 Antiviral Activity of 2-Thio-6-Azauridine vs YF Virus: A summary of the antiviral and cytotoxicity performance of the second control compound, AVS-6724 (2-Thio-6-Azauridine) is presented in Figure 51-B for 52 tests performed during June 1989 through January, 1991.

Second Control Compound-Antiviral Performance: 2-Thio-6-Azauridine (AVS-6724) has been tested as a possible second control compound against YF in these MTT-assay prescreens. The mean and median antiviral inhibition and cytotoxicity patterns of the second positive control drug are illustrated in Figure 51-B.

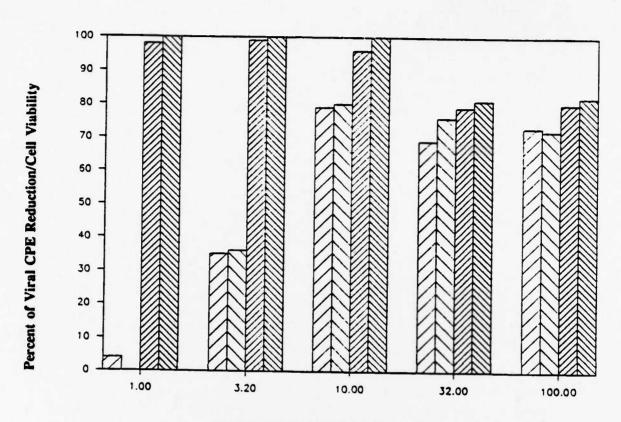
The 52 control tests performed with 2-Thio-6-Azauridine gave a mean Selectivity Index (SI) of 8.14 (SD \pm 4.60) and the median value was 7.70, indicating moderate antiviral selectivity for 2-Thio-6-Azauridine. The reason for this discrepancy is that even at 100 μ g/ml the maximum cytotoxic effect does not consistently reach 25% reduction level, thus SI calculations cannot be executed properly. (SI is calculated by dividing the TC₂₅ by the IC₅₀).

The mean Antiviral Inhibitory Concentration 50% (IC₅₀) was 5.66 μ g/ml (SD \pm 4.75). The median IC₅₀ value was 4.74 μ g/ml. This indicates that 2-Thio-6-Azauridine does reach 50% antiviral reduction levels, consistently at 10 μ g/ml concentration. During this reporting period, the highest concentration of 2-Thio-6-Azauridine was varied from 1 to 320 μ g/ml to properly evaluate the maximum antiviral effect and cytotoxicity pattern of 2-Thio-6-Azauridine.

The maximum antiviral inhibitory level of 52 2-Thio-6-Azauridine tests (Figure 51-B) varies depending upon which drug concentration scale is used. Maximum antiviral effect ($\sim 80\%$) was found with a simultaneous $\sim 5\%$ cytotoxic suppression. Above 10 μ g/ml concentration 2-Thio-6-Azauridine starts to lose its antiviral potency with increasing cytotoxicity. An increase of the concentration of 2-Thio-6-Azauridine to 100 μ g/ml does not improve the antiviral activity (Figure 51-B).

In the present 52 assays we tested 2-Thio-6-Azauridine at 0.5 \log_{10} scale concentrations (1 - 32 μ g/ml) to maximize the correct measurements of its antiviral and cytotoxicity effects. This enabled us to monitor our quality control parameters more accurately.

2-THIO-6-AZAURIDINE -VS- YF (PRE-SCREEN) VIRUS



CONCENTRATION (µg/ml)

Wiral CPE
Reduction

Median % Viral CPE Reduction

Mean % Cell Viability

Median % Cell Viability

% Viral CPE Reduction

% Cell Viability

Conc.(µg/ml)	1	3.2	10	32	100	1	3.2	10	32	100
Mean	4	35	79	69	73	98	99	96	79	80
Median	0	36	80	76	72	100	100	100	81	82
Std. Dev.	0.09	0.24	0.19	0.23	0.08	0.04	0.03	0.06	0.14	0.08

Figure 51-B

Average Antiviral and Cytotoxicity Values for 52 Positive Control Compound Tests

4.2.1.1.2 <u>Maximum Antiviral Effect of 2-Thio-6-Azauridine vs YF Virus:</u> Since the metabolic activity of the cells was an unknown function during the testing period, it was monitored indirectly by measuring the maximum antiviral effect of the control compound 2-Thio-6-Azauridine. This demonstrated the amount of infectious virus produced by the cells (Maximum Percent CPE).

A bar graph scatter plot (Figure 52-B) depicts the distribution of the maximum antiviral reduction values of all 52 control compound prescreen assays for 2-Thio-6-Azauridine. The results indicate that the average maximum antiviral reduction obtained with the present SOP is around 86% (SD \pm 13.50) reduction levels. The maximum reduction levels vary from 56 - 100% but remain quite consistently around the median of 87.5%. The assay control values give a shifted bell-shaped distribution curve toward the median 87.5% reduction level. This indicates quite a consistent day-to-day performance of the control compound in the YF prescreen-MTT assay.

During this period the positive control compound performance criteria for 2-Thio-6-Azauridine versus the YF virus in the pre-screen format has not been set to a definite endpoint. We have collected data in order to find out what would constitute a reasonable quality control endpoint and the data indicates that it could be set at the 50% reduction level. In order to measure the maximum antiviral endpoints of 2-Thio-6-Azauridine correctly, the concentration scale and the highest concentration to be used, must be evaluated at the proper (semi-log) scale $(1 - 32 \mu g/ml)$ as seen in Figure 51-B.

2-Thio-6-Azauridine is active in vitro against YF virus and functions as a better quality control compound than our present control, Selenazofurin.

Variation of the Maximum Antiviral Effect YF Virus - VS - 2-Thio-6-Azauridine (Prescreen Protocol)

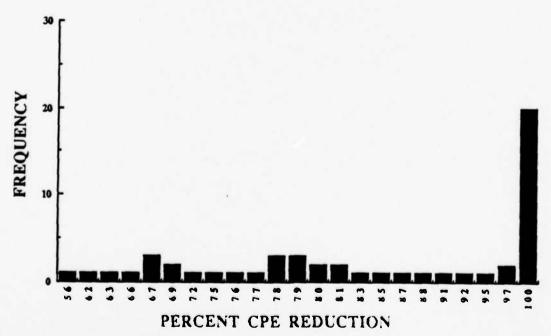


Figure 52-B

Maximum Antiviral CPE Reduction (%).

Summary of 52 Control Tests.

4.2.1.1.3 Cellular Cytotoxicity of 2-Thio-6-Azauridine vs YF Virus:

<u>YF-Control Compound-Cytotoxicity Performance</u>: The 52 cytotoxicity values of the positive control compound 2-Thio-6-Azauridine are also very consistent. The mean cell Toxic Concentration 25% (TC₂₅) was 35.5 μ g/ml (SD \pm 22.00) and the median was 32.00 μ g/ml (range of 9.75 - 108 μ g/ml).

As can be seen from Figure 51-B, no definite TC_{25} toxicity value can be consistently measured with a 100 μ g/ml log_{10} scale. Further increase in the concentration of 2-Thio-6-Azauridine would be needed to properly evaluate the maximum cytotoxicity of 2-Thio-6-Azauridine.

Figure 51-B, indicates that when the cytotoxicity reaches ~ 0 - 5% at 10 μ g/ml, the control compound (2-Thio-6-Azauridine) has reached simultaneously its maximum antiviral effect. The cytotoxic effect of 2-Thio-6-Azauridine is insignificant between 1 and 10 μ g/ml. The average cytotoxicity reached $\sim 20\%$ at 100 μ g/ml, which was the highest 2-Thio-6-Azauridine concentration in most tests.

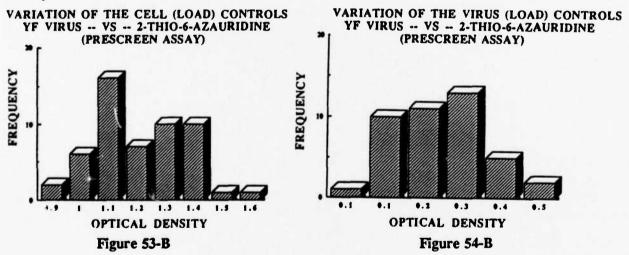
2-Thio-6-Azauridine has a definite cytotoxic suppression on cellular metabolism and growth. However, the TC_{25} and TC_{50} toxicity could not be consistently achieved with the 100 μ g/ml concentration of 2-Thio-6-Azauridine. Therefore, a readjustment to 320 μ g/ml (as being the highest 2-Thio-6-Azauridine concentration tested) would be needed to properly evaluate the TC_{25} endpoint. However, at this concentration (100 μ g/ml) the IC_{50} cannot be measured consistently.

4.2.1.1.4 YF-Assay Plate Ouality Controls: Cell Load and Virus Load Parameters (2-Thio-6-Azauridine): The MTT assay is fundamentally dependent upon the quality of the assay plates. Our large-scale antiviral testing is dependent upon the uniformity of the test plates produced for the daily assays. Equal loads of cell load and virus load as well as the consistent performance of the reagents used daily was monitored. A sample of the plate variation control for the period of June, 1989 through January, 1991 is presented in Figures 53-B, 54-B, and 55-B.

<u>YF-Control Compound-Cell Load Performance</u>: A bar graph scatter plot of the mean cell control (O.D. reading) of 52 control assays is plotted in Figure 53-B. The results indicate that the cell O.D. readings reached a mean 1.210 (SD \pm 0.160) with a median of 1.210 (range of 0.890 - 1.570). This indicates that a uniform and equal number (18,000 cells/well) of cells are being loaded into every well in the 96-well plate during the day-to-day operation. The cells reduced MTT to formazan giving maximum blue color uniformly and consistently.

<u>YF-Control Compound-Virus Load Performance</u>: A bar graph scatter plot of the mean virus load O.D. readings of the 52 control assays is presented in Figure 54-B. The results indicate that the average virus load O.D. reading is 0.350 (SD \pm 0.140) with a median of 0.340 (range of 0.140 - 0.680). This demonstrates that a reasonable cell destruction is taking place and a uniform load of virus (32 TCID₅₀) is administered on the cell monolayer with consistent viral CPE results.

YF-Control Compound-Assay Differential Performance: A bar graph scatter plot of the mean O.D. differential values of the 52 control assays is provided in Figure 55-B. The results indicate that the average differential O.D. reading is 0.850 (SD \pm 0.130) with a median of 0.860 (range 0.538 - 1.195). The single bell-shaped curve is reasonably sharp and uniform. This reflects that the assays are executed consistently and are repeatable during day-to-day operation with close to 86% measurement accuracy.



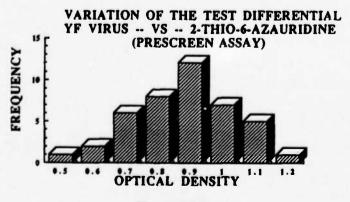


Figure 55-B

4.2.1.2 Prescreen YF-Antiviral Activity Results:

New Drugs with 50% Antiviral Reduction Levels: Out of the 3606 accepted single drug tests, 61 new compounds demonstrated antiviral activity, having antiviral reduction values better than 50%. This represents around 1.7% of the test compounds being active at this antiviral reduction level. These compounds are summarized in Table 31 according to the highest Selectivity Index (SI). One compound, B723123, demonstrated the best antiviral promise having an SI of 325. Six compounds demonstrated moderate antiviral activity, having SI's from 22 - 56. The rest (54) demonstrated minimal antiviral selectivity with SI's that ranged from <1 - 11.

Table 31

New Prescreen Drugs that Produced 50% Antiviral Reduction Against YF Virus

DRUG	VIR	PLT	SHIP	TEST		WC 25		TO 50		0.7
	ATK	•	•	DATE		TC 25		IC 50		SI
B723123	YF	X2V	5P	06/20/90		325.00	<	1.00	>	325.00
B721173	YF	VOO	4P	04/05/90		325.00		5.80		56.05
B722182	YF	OK8		11/22/89		163.00		3.70		44.17
B722116	YF	ZDQ	9P	10/09/90		182.00		4.43		41.10
B722241	YF	OKD		11/22/89	>			31.60	>	31.62
B723412	YF	XIO	6P	07/12/90		227.00		7.99		28.40
B721568	YF	OAE		08/09/89		360.00		16.20		22.24
4279	YF	09W		08/03/89		32.50		2.97		10.95
4280	YF	09W		08/03/89		32.50		3.00		10.83
B721511	YF	09X		08/03/89		330.00		30.50		10.81
B721899	YF	0J2		11/08/89		325.00		30.90		10.52
B724898	YF	OFN		09/27/89		322.00		37.30		8.65
B722183	YF	OK8		11/22/89		270.00		36.10		7.48
GRP19380	YF	WO4	4P	06/07/90		10.40		1.40		7.46
B724618	YF	OAN		08/09/89		248.00		37.10		6.69
B721823	YF	OGC		10/04/89	>	1000.0		154.00	>	6.51
4276	YF	09V		08/03/89		32.50		5.06		6.42
B724610	YF	OAL		08/09/89		262.00		45.10		5.82
B722239	YF	OKD		11/22/89		244.00		43.60		5.60
B724701	YF	OCQ		08/24/89		227.00		42.20		5.39
B724592	YF	OAI		08/09/89		204.00		41.40		4.94
B724466	YF	07Y		07/12/89		218.00		45.70		4.77
2630	YF	05N		06/21/89		100.00		22.30		4.49
B722111	YF	OJQ		11/15/89		175.00		45.30		3.85
B721754	YF	065		06/29/89		360.00		97.10		3.71
GRP19396	YF	WOK	4P	06/08/90		230.00		62.20		3.70
B724468	YF	07Y		07/12/89		329.00		98.80		3.33
B723141	YF	X2Y	5P	06/20/90				316.00		3.16
B721953	YF	0J2		11/08/89				320.00		3.13
B722165	YF	OJV		11/15/89	>			345.00	>	2.90
B721728	YF	05Q		06/21/89		1000.0		447.00		2.24
B724844	YF	OEZ		09/14/89		110.00		50.50		2.18
B724852	YF	OF1		09/14/89		142.00		67.90		2.09
B722246	YF	OKE		11/22/89		118.00		62.70		1.89
B724530	YF	09G		07/26/89		126.00		68.70		1.84
B724863	YF	OFG		09/27/89		125.00		69.30		1.81
B721823	YF	013		10/25/89	>			574.00	>	1.74
B722109	YF	OJQ		11/15/89		155.00		90.70		1.71
B724812	YF	OET		09/14/89		135.00		80.10		1.69
B722824	YF	UBS		03/07/90	_	53.10		31.50		1.68
B722248	YF	151		11/08/90	>			602.00	>	1.66
B722518	YF		3P	03/16/90		942.00		586.00		1.61
B721702	YF	05M		06/21/89		1000.0		748.00		1.34
B722181	YF	OJW		11/15/89		123.00		93.90		1.31
B721166	YF	09V		08/03/89		589.00		451.00		1.30
B724772	YF	OG4	10	10/04/89		92.80		73.90		1.26
B723441	YF	ZRY		10/30/90				805.00		1.24
B723427	YF	ZRU	10	10/26/90	>			890.00	>	1.12
B721714	YF	050		06/21/89		1000.0		919.00		1.09
B722247	YF	OKE		11/22/89		69.70		65.50		1.06

Table 31 (Cont'd)

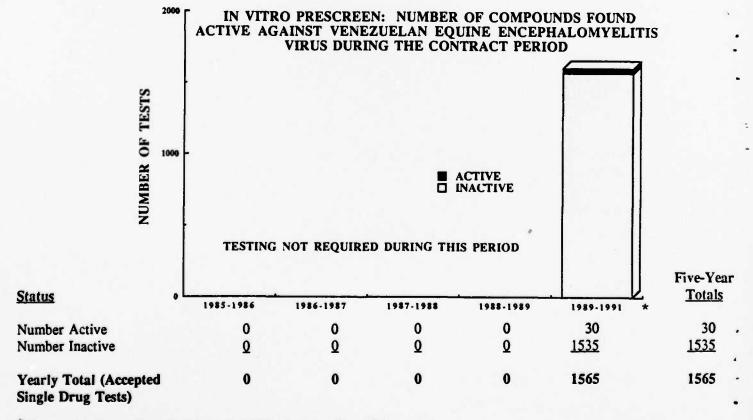
DRUG VII		SHIP	TEST DATE		TC 25	IC 50		SI
*			DALE		10 23	10 30		31
B722087 YF	OJA		11/08/89		736.00	703.00		1.05
B724442 YF	07T		07/12/89		730.00	715.00		1.02
B723436 YF	ZRX	10	10/30/90	>	1000.0	1000.0	>	1.00
B848752 YF	2M6	11P	01/17/91	>	3200.0	3200.0	>	1.00
B724654 YF	OBH		08/16/89		8.08	8.54		0.95
B724510 YF	09C		07/26/89		698.00	808.00		0.86
B724731 YF	0D6		08/30/89		598.00	692.00		0.86
B724663 YF	OBJ		08/16/89		56.80	68.10		0.83
B724657 YF	OBI		08/16/89		493.00	786.00		0.63
B721559 YF	09V		08/03/89		457.00	798.00		0.57
B721770 YF	067		06/29/89		79.10	316.00		0.25

The *in vitro* antiviral activity of the compounds in Table 31 should be further confirmed. Verification of the antiviral activity of these prescreen actives was done using the primary screening (confirmatory) protocol. (See Table 34)

4.2.2 Prescreen Assay (Venezuelan Equine Encephalomyelitis Virus [VEI):

The number of single drug tests carried against VE during this contract period is summarized in yearly increments in Figure 56. During this seven-month period (June, 1990 - January, 1991) 2235 tests were performed against the VE-virus with the MTT-assay format. Out of these, 126 were control compound assays-selenazofurin (AVS-0253) being the primary positive control compound. Three hundred ninety-nine (399) tests were internal (+++) virus load, cell load, and other quality control tests. One hundred forty-five (145) tests were considered unsatisfactory based on the preliminary criteria of the quality controls set during this reporting period. The rest, totaling 1565 were actual single drug tests. The total number of tests (2235) represents testing for an seven-month period since we only added the VE virus to the prescreen assay protocol in June 1990. The 145 unsatisfactory tests represent a 6.5% rejection rate based on present preliminary quality control parameters used for the VE-virus.

Out of the 1565 test compounds, 30 demonstrated antiviral activity at greater than 50% reduction levels. This represents 2% of the tested compounds having *in vitro* antiviral activity against VE-virus. The remainder, 1535 compounds (98%), are to be considered inactive with present quality control and assay protocols.



Represents 7-month period (June 1, 1990 - January 31, 1991)

Figure 56

- 4.2.2.1 Prescreen VE-Quality Controls: Two positive control compounds (Selenazofurin and 2-Thio-6 Azauridine) were used in the daily assay sets as antiviral activity quality controls. The antiviral performance of the unknown compounds is compared to that of the positive control compounds. Compounds with equal to or better antiviral potency are considered active and are worthy of further in vitro profile studies and in vivo testing.
- 4.2.2.1.1 <u>Antiviral Activity of Selenazofurin vs VE Virus:</u> A summary of the antiviral and cytotoxicity performance of the primary control compound, AVS-0253 (Selenazofurin) is presented in Figure 57-A for 119 tests performed during June, 1990 through January, 1991.

Control Compound-Antiviral Performance: Selenazofurin (AVS-0253) has been the sole control compound against VE in these MTT-assay prescreens. The mean and median antiviral inhibition a cytotoxicity patterns of the positive control drug (Selenazofurin) are illustrated in Figure 57-A.

The 119 control tests performed with Selenazofurin gave a mean Selectivity Index (SI) of 4.60 (SD \pm 6.60) and the median value was 2.40, indicating poor antiviral selectivity for Selenazofurin. SI is calculated by dividing the TC_{25} by the IC_{50} .

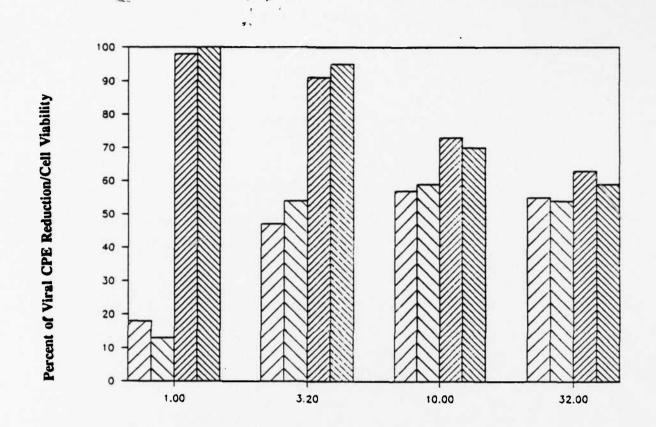
The mean Antiviral Inhil pry Concentration 50% (IC₅₀) was 5.30 μ g/ml (SD \pm 7.40). The median IC₅₀ value was 2.60 μ g/ml. This indicates that Selenazofurin does reach 50% antiviral reduction levels reasonably well at 3.2 - 32 μ g/ml concentrations (Figure 57-A). During this reporting period, the highest and lowest concentrations of Selenazofurin was varied from 0.32 to 320 μ g/ml to properly evaluate the maximum antiviral effect and cytoto..icity pattern of Selenazofurin. These statistics are based on 119 values tested using the 1 - 32 μ g/ml 0.5 \log_{10} scale.

The average maximum antiviral inhibitory level of 119 Selenazofurin tests (Figures 57-A) was reached at 10 μ g/ml of compound with 59% antiviral effect. Maximum antiviral effect (~59%) was found with a simultaneous ~30% cytotoxic suppression. Above this concentration (10 μ g/ml) Selenazofurin starts to lose its antiviral potency with increasing cytotoxicity (Figure 57-A). An increase of the concentration of Selenazofurin to 100 - 320 μ g/ml does not improve the antiviral activity (Figure 57-A). Actually antiviral activity decreases from 59% (at 10 μ g/ml) to ~10% (at 320 μ g/ml).

As reported previously in the 9th semiannual report, depending upon which concentration scale is used (log or semilog scale) a different maximum antiviral value is obtained.

In the present, we tested 119 control assays with Selenazofurin at 0.5 \log_{10} scale from 1 - 32 μ g/ml to maximize the correct measurements of its antiviral and cytotoxicity effects. This enabled us to monitor our quality control parameters more accurately.

SELENAZOFURIN -VS- VE VIRUS (PRESCREEN PROTOCOL)



CONCENTRATION (µg/ml)

Reduction Reduction	Wean % Viral CPE Reduction	Median % Viral CPE Reduction	Cell Viability	Median % Cell Viabili
---------------------	----------------------------	------------------------------	----------------	-----------------------

% Viral CPE	Reduction	% Cell	Viability

Conc. (µg/ml)	1	3.2	10	32	1	3.2	10	32
Mean	18	47	57	55	98	91	73	63
Median	13	54	59	54	100	95	70	59
Std. Dev.	0.16	0.26	0.21	0.14	.04	0.11	0.18	0.18

Figure 57-A

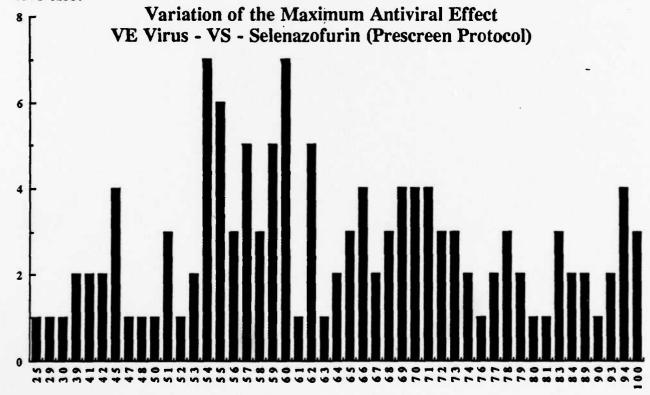
Average Antiviral and Cytotoxicity Values for 119 Positive Control Compound Assays

4.2.2.1.2 <u>Maximum Antiviral Effect of Selenazofurin vs VE Virus</u>: Since the metabolic activity of the cells was an unknown function during the testing period, it was monitored indirectly by measuring the maximum antiviral effect of the control compound Selenazofurin. This demonstrated the amount of infectious virus that was produced by the cells (Maximum Percent CPE).

A bar graph scatter plot (Figure 58-A) depicts the distribution of the maximum antiviral reduction values of all 119 control compound prescreen assays for Selenazofurin. The results indicate that the average maximum antiviral reduction obtained with the present SOP is around 64% (SD \pm 15.20) reduction levels. The maximum reduction levels vary from 25 - 100% but remain quite consistently around the median of 62%. The assay control values give a reasonable bell-shaped distribution curve toward the median 62% reduction level. This indicates quite a consistent day-to-day performance of the control compound in the VE prescreen-MTT assay.

During this period the positive control compound performance criteria for Selenazofurin versus the VE virus in the pre-screen format has not been set to a definite endpoint. We have collected data in order to find out what would constitute a reasonable quality control endpoint and the data indicates that it could be set at the 50% reduction level. In order to measure the maximum antiviral endpoints of Selenazofurin correctly, the concentration scale and the highest concentration to be used, must be evaluated at the proper (semi-log) scale as seen in Figure 57-A.

Selenazofurin is active *in vitro* against VE virus and functions as a reasonable quality control compound. On the other hand, regardless of the performance of the VE-quality control drug Selenazofurin, around 30 other compounds have equal or better antiviral activity against VE virus than AVS-0253.



FREQUENCY

PERCENT CPE REDUCTION

Figure 58-A
Maximum Antiviral CPE Reduction (%).
Summary of 119 Control Tests.

4.2.2.1.3 Cellular Cytotoxicity of Selenazofurin vs VE Virus:

<u>VE-Control Compound-Cytotoxicity Performance:</u> The 119 cytotoxicity values of the positive control compound Selenazofurin are also very consistent. The mean cell **Toxic Concentration 25%** (TC₂₅) was 14.40 μ g/ml (SD \pm 11.10) and the median was 8.40 μ g/ml (range of 1.0 - 32 μ g/ml). The reason for this discrepancy is that at 32 μ g/ml scale, neither the TC₂₅ cytotoxicity or the IC₅₀ cannot always be measured accurately.

As can be seen from Figure 57-A, a definite TC_{25} toxicity value can be measured with a 32 μ g/ml \log_{10} scale. Further increase in the concentration of Selenazofurin would be needed to consistently evaluate the maximum cytotoxicity of Selenazofurin. Also, Figure 57-A indicates that when the cytotoxicity reaches ~25 - 30% at 10 μ g/ml, the control compound (Selenazofurin) has reached simultaneously its maximum antiviral effect. The cytotoxic effect of Selenazofurin is insignificant below 3.2 μ g/ml. The average cytotoxicity reached ~30% at 32 μ g/ml, which was the highest Selenazofurin concentration in most tests.

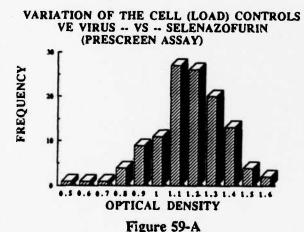
Selenazofurin has a definite cytotoxic suppression on cellular metabolism and growth, however, the TC_{25} and TC_{50} toxicity can be achieved with reasonable consistency at 32 μ g/ml concentration. Therefore, a readjustment to 100 μ g/ml (as being the highest Selenazofurin concentration tested) is not needed. However, at this concentration (100 μ g/ml) the IC₅₀ cannot yet be measured accurately.

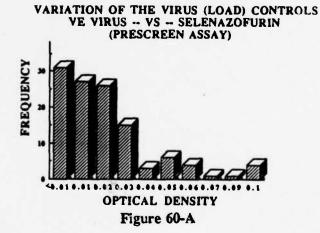
4.2.2.1.4 <u>YE-Assay Plate Ouality Controls: Cell Load and Virus Load Parameters:</u> The MTT assay is fundamentally dependent upon the quality of the assay plates. Our large-scale antiviral testing is dependent upon the uniformity of the test plates produced for the daily assays. Equal numbers of cell load and virus load as well as the consistent performance of the reagents used daily was monitored. A sample of the plate variation control for the period of June, 1990 through January, 1991, is presented in Figures 59-A, 60-A and 61-A.

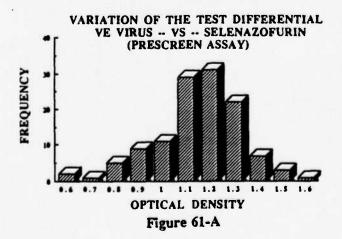
<u>VE-Control Compound-Cell Load Performance</u>: A bar graph scatter plot of the mean cell control (O.D. reading) of 119 control assays is plotted in Figure 59-A. The results indicate that the cell O.D. readings reached a mean 1.200 (SD \pm 0.150) with a median of 1.230 (range of 0.530 - 1.600). This indicates that a uniform and equal number (18,000 cells/well) of cells are being loaded into every well in the 96-well plate during the day-to-day operation. The cells reduced MTT to formazan giving maximum blue color uniformly and consistently.

<u>VE-Control Compound-Virus Load Performance</u>: A bar graph scatter plot of the mean virus load O.D. readings of the 119 control assays is presented in Figure 60-A. The results indicate that the average virus load O.D. reading is 0.020 (SD \pm 0.030) with a median of 0.020 (range of 0 - 0.160). This demonstrates that a reasonable cell destruction is taking place and a uniform load of virus (32 TCID₅₀) is administered on the cell monolayer with consistent viral CPE results.

<u>VE-Control Compound-Assay Differential Performance:</u> A bar graph scatter plot of the mean O.D. differential values of the 119 control assays is provided in Figure 61-A. The results indicate that the average differential O.D. reading is 1.140 (SD \pm 0.180) with a median of 1.160 (range 0.572 - 1.568). The single bell-shaped curve is reasonably sharp and uniform. This reflects that the assays are executed consistently and are repeatable during day-to-day operation with close to 85% measurement accuracy.







4.2.2.1 Prescreen VE-Quality Controls:

4.2.2.1.1 Antiviral Activity of 2-Thio-6-Azauridine vs VE Virus: A summary of the antiviral and cytotoxicity performance of the second control compound, AVS-6724 (2-Thio-6-Azauridine) is presented in Figure 57-B for 50 tests performed during June, 1990 through January, 1991.

<u>Second Control Compound-Antiviral Performance</u>: 2-Thio-6-Azarridine (AVS-6724) has been tested as a possible second control compound against VE in these MTT-assay prescreens. The mean and median antiviral inhibition and cytotoxicity patterns of the second positive control drug are illustrated in Figure 57-B.

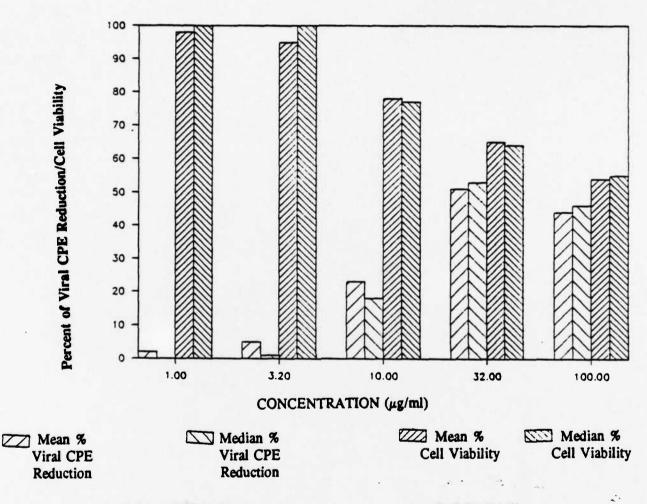
The 50 control tests performed with 2-Thio-6-Azauridine gave a mean Selectivity Index (SI) of 0.69 (SD \pm 0.78) and the median value was 0.42, indicating poor antiviral selectivity for 2-Thio-6-Azauridine. The reason for this discrepancy is that even at 100 μ g/ml the maximum antiviral effect does not consistently reach 50% reduction level, thus SI calculations cannot be executed properly. SI is calculated by dividing the TC₂₅ by the IC₅₀.

The mean Antiviral Inhibitory Concentration 50% (IC₅₀) was 13.00 μ g/ml (SD \pm 12.00). The median IC₅₀ value was 13.70 μ g/ml. This indicates that 2-Thio-6-Azauridine does reach 50% antiviral reduction levels, with reasonable consistently at 32 μ g/ml maximum concentration. During this reporting period, the highest and lowest concentration of 2-Thio-6-Azauridine was varied from 1 to 320 μ g/ml to properly evaluate the maximum antiviral effect and cytotoxicity pattern of 2-Thio-6-Azauridine. The statistics are based on 50 assays tested using the 1 - 100 μ g/ml 0.5 log₁₀ scale.

The average maximum antiviral inhibitory level of 50 2-Thio-6-Azauridine tests (Figure 57-B) was reached at 32 μ g/ml of the compound with 53% antiviral effect. Maximum antiviral effect (~53%) was found with a simultaneous ~35% cytotoxic suppression. Above 32.0 μ g/ml concentration 2-Thio-6-Azauridine starts to lose its antiviral potency with increasing cytotoxicity. An increase of the concentration of 2-Thio-6-Azauridine to 100 - 320 μ g/ml does not improve the antiviral activity (Figure 57-B). Actually antiviral activity decreases from 53% (at 32 μ g/ml) to ~10% (at 320 μ g/ml).

In the present 50 assays, we tested 2-Thio-6-Azauridine at 0.5 log₁₀ scale concentrations to maximize the correct measurements of its antiviral and cytotoxicity effects. This enabled us to monitor our quality control parameters more accurately.

2-THIO-6-AZAURIDINE -VS- VE (PRE-SCREEN) VIRUS



% Viral CPE Reduction

% Cell Viability

Conc.(µg/ml)	1	3.2	10	32	100	1	3.2	10	32	100
Mean	2	5	23	51	44	98	95	78	65	54
Median	0	1	18	53	46	100	100	77	64	55
Std. Dev.	0.07	0.09	0.21	0.13	0.11	0.05	0.09	0.15	0.10	0.11

Figure 57-B

Average Antiviral and Cytotoxicity Values for 50 Positive Control Compound Tests

4.2.2.1.2 <u>Maximum Antiviral Effect of 2-Thio-6-Azauridine vs VE Virus:</u> Since the metabolic activity of the cells was an unknown function during the testing period, it was monitored indirectly by measuring the maximum antiviral effect of the control compound 2-Thio-6-Azauridine. This demonstrated the amount of infectious virus produced by the cells (Maximum Percent CPE).

A bar graph scatter plot (Figure 58-B) depicts the distribution of the maximum antiviral reduction values of all 50 control compound prescreen assays for 2-Thio-6-Azauridine. The results indicate that the average maximum antiviral reduction obtained with the present SOP is around 52% (SD \pm 13.50) reduction levels. The maximum reduction levels vary from 23 - 80% but remain quite consistently around the median of 54%. The assay control values give a reasonable bell-shaped distribution curve toward the median 54% reduction level. This indicates quite a consistent day-to-day performance of the control compound in the VE prescreen-MTT assay.

During this period the positive control compound performance criteria for 2-Thio-6-Azauridine versus the VE virus in the pre-screen format has not been set to a definite endpoint. We have collected data in order to find out what would constitute a reasonable quality control endpoint and the data indicates that it could be set at the 25% reduction level. In order to measure the maximum antiviral endpoints of 2-Thio-6-Azauridine correctly, the concentration scale and the highest concentration to be used, must be evaluated at the proper (semi-log) scale of $1 - 32 \mu g/ml$ as seen in Figure 57-B.

2-Thio-6-Azauridine is active in vitro against VE virus and functions as a reasonable quality control compound similar to the present control compound Selenazofurin.

Variation of the Maximum Antiviral Effect VE Virus - VS - 2-Thio-6-Azauridine (Prescreen Protocol)

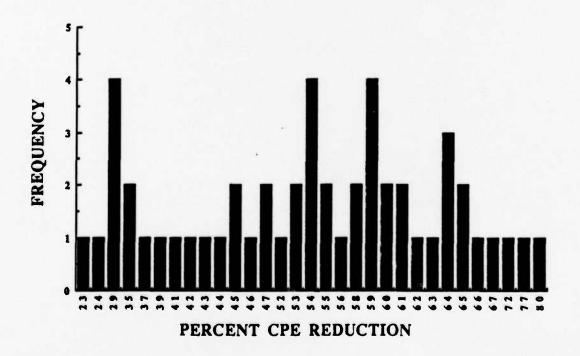


Figure 58-B

Maximum Antiviral CPE Reduction (%).

Summary of 50 Control Tests.

4.2.2.1.3 Cellular Cytotoxicity of 2-Thio-6-Azauridine vs VE Virus:

<u>VE-Control Compound-Cytotoxicity Performance</u>: The 50 cytotoxicity values of the positive control compound 2-Thio-6-Azauridine are also very consistent. The mean cell **Toxic Concentration** 25% (TC₂₅) was 18.70 μ g/ml (SD \pm 14.70) and the median was 13.00 μ g/ml (range of 1.02 - 83.1 μ g/ml).

As can be seen from Figure 57-B, a definite TC_{25} toxicity value can be consistently measured with 32 μ g/ml at 0.5 log_{10} scale. Further increase in the concentration of 2-Thio-6-Azauridine would be needed to properly evaluate the maximum cytotoxicity.

Figure 57-B indicates that when the cytotoxicity reaches ~ 25 - 30% at 32 μ g/ml, the control compound (2-Thio-6-Azauridine) as reached simultaneously its maximum antiviral effect. The cytotoxic effect of 2-Thio-6-Azauridine is insignificant below 3.2 μ g/ml. The average cytotoxicity reached $\sim 45\%$ at 100 - 320 μ g/ml, which was the highest 2-Thio-6-Azauridine concentration in most tests.

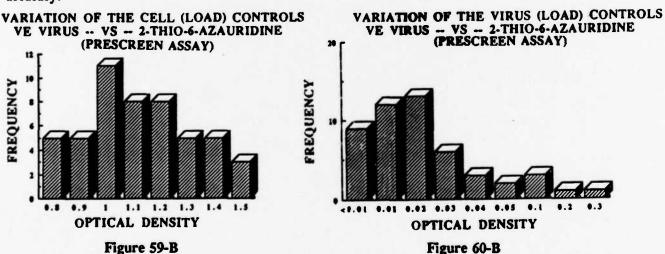
2-Thio-6-Azauridine has a definite cytotoxic suppression on cellular metabolism and growth, however, the TC_{25} and TC_{50} toxicity cannot be consistently measured at 100 μ g/ml concentration of 2-Thio-6-Azauridine. Therefore, a readjustment to 100 - 320 μ g/ml (as being the highest 2-Thio-6-Azauridine concentration tested) would be needed to properly evaluate the TC_{25} and TC_{50} endpoints. However, at this concentration (100 μ g/ml) the IC_{50} cannot yet be measured accurately.

4.2.2.1.4 <u>VE-Assay Plate Quality Controls: Cell Load and Virus Load Parameters (2-Thio-6-Azauridine):</u> The MTT assay is fundamentally dependent upon the quality of the assay plates. Our large-scale antiviral testing is dependent upon the uniformity of the test plates produced for the daily assays. Equal loads of cell load and virus load as well as the consistent performance of the reagents used daily was monitored. A sample of the plate variation control for the period of June, 1990 through January, 1991 is presented in Figures 59-B, 60-B, and 61-B.

<u>VE-Control Compound-Cell Load Performance</u>: A bar graph scatter plot of the mean cell control (O.D. reading) of 50 control assays is plotted in Figure 59-B. The results indicate that the cell O.D. readings reached a mean 1.120 (SD \pm 0.190) with a median of 1.120 (range of 0.810 - 1.550). This indicates that a uniform and equal number (18,000 cells/well) of cells are being loaded into every well in the 96-well plate during the day-to-day operation. The cells reduced MTT to formazan giving maximum blue color uniformly and consistently.

<u>VE-Control Compound-Virus Load Performance</u>: A bar graph scatter plot of the mean virus load O.D. readings of the 50 control assays is presented in Figure 60-B. The results indicate that the average virus load O.D. reading is 0.030 (SD \pm 0.060) with a median of 0.020 (range of 0 - 0.330). This demonstrates that a reasonable cell destruction is taking place and a uniform load of virus (32 TCID₅₀) is administered on the cell monolayer with consistent viral CPE results.

<u>VE-Control Compound-Assay Differential Performance:</u> A bar graph scatter plot of the mean O.D. differential values of the 50 control assays is provided in Figure 61-B. The results indicate that the average differential O.D. reading is 1.090 (SD \pm 0.190) with a median of 1.080 (range 0.786 - 1.519). The single bell-shaped curve is reasonably sharp and uniform. This reflects that the assays are executed consistently and are repeatable during day-to-day operation with close to 92% measurement accuracy.



VARIATION OF THE TEST DIFFERENTIAL
VE VIRUS -- VS -- 2-THIO-4-AZAURIDINE
(PRESCREEN ASSAY)

FREQUENCY

OPTICAL DENSITY
Figure 61-B

4.2.2.2 Prescreen VE-Antiviral Activity Results:

New Drugs with 50% Antiviral Reduction Levels: Out of the 1565 accepted single drug tests, 30 new compounds demonstrated antiviral activity, having antiviral reduction values better than 50%. This represents around 2.0% of the test compounds being active at this antiviral reduction level. These compounds are summarized in Table 32 according to the highest Selectivity Index (SI). B723096 demonstrated the best antiviral promise having a SI of around 5.0. Three other compounds demonstrated moderate antiviral activity, having SI's that ranged from 3 - 4. The other 26 compounds showed some degree of activity having SI's that ranged from 0.4 - 2.4.

Table 32

New Prescreen Drugs that Produced 50% Antiviral Reduction Against VE Virus

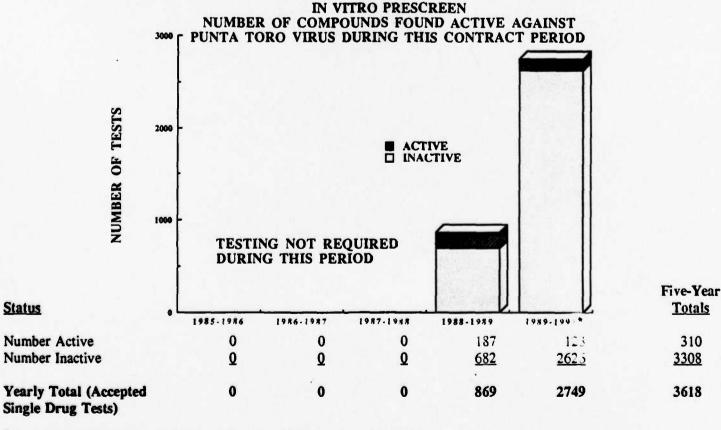
DRUG		PLT	SHIP	TEST				
*	VIR	#	#	DATE		TC 25	IC 50	SI
B723096	VE	WWY	5P	06/19/90	>	1000.0	208.00 >	4.80
B720933	VE	Z15	8P	09/28/90	>	1000.0	277.00 >	3.62
B723322	VE	XIC	6P	07/13/90	>	1000.0	343.00 >	2.91
B723148	VE	WWT	5P	06/19/90	>	1000.0	382.00 >	2.62
B723414	VE	UHX	6P	07/13/90			419.00 >	2.39
B723802	VE	ZTP	10	11/02/90	>	1000.0	425.00 >	2.35
B723315	VE	XSE	6P	07/27/90	>	1000.0	466.00 >	2.15
B848727	VE	2J6	11P	01/17/91		3200.0	1490.0 >	2.15
B723247	VE	XHR	5P	07/13/90	>	1000.0	488.00 >	2.05
B723103	VE	WWZ	5P	06/19/90		1000.0	492.00 >	2.03
B848653	VE	2J4	11P	01/17/91		2670.0	1350.0	1.97
B723061	VE	WWW	5P	06/19/90	>	1000.0	525.00 >	1.91
B723250	VE	XO9	6P	07/20/90	>	1000.0	580.00 >	1.73
B723465	VE	ZSA	10	10/30/90		981.00	569.00	1.72
B719218	VE	220	12P	12/18/90	>	1000.0	589.00 >	1.70
B723141	VE	WWS	5P	06/19/90		765.00	453.00	1.69
B723431	VE	14H	10P	11/09/90		880.00	537.00	1.64
B723409	VE	XHU	6P	07/13/90	>	1000.0	688.00 >	1.45
B723203	VE	X7W	5P	06/26/90		902.00	656.00	1.37
B723807	VE	ZTQ	10	11/02/90		720.00	596.00	1.21
B723136	VE	WWR		06/19/90	>	1000.0	890.00 >	1.12
B723318	VE	XSF	6P	07/27/90	>	1000.0	1000.0 >	1.00
B723240	VE	X8N		06/29/90		628.00	680.00	0.92
B723364	VE	XJA		07/17/90		704.00	761.00	0.92
B723140	VE	WWR		06/19/90		823.00	907.00	0.91
B848911	VE		12P	12/11/90		685.00	825.00	0.83
B723232	VE	X8L		06/29/90		557.00	710.00	0.78
B723391	VE	X07	6P	07/20/90		474.00	861.00	0.55
B723064	VE	WWW		06/19/90		508.00	1000.0	0.51
B849177	VE	XIB		07/13/90		278.00	746.00	0.37
				,, 50				

The *in vitro* antiviral activity of the compounds in Table 32 was further confirmed in most of the compounds. Verification of the antiviral activity of these prescreen actives was tested using the primary screening (confirmatory) protocol.

4.2.3 Prescreen Assay (Punta Toro Virus [PT])

The number of single drug tests carried against PT during this contract period is summarized in yearly increments in Figure 62. During this nineteen-month period (June, 1989 - January, 1991) 6026 tests were performed against the PT-virus with MTT-assay format. Out of these, 298 were control compound assays-ribavirin (AVS-0001) being the primary positive control compound. Nine hundred seventy-six (976) tests were internal (+++) virus load, cell load, and other quality control tests. One thousand one hundred thirty-four (1134) tests were considered unsatisfactory based on the preliminary criteria of the quality controls set during this reporting period. The rest, totaling 3618 were actual single drug tests. The 1134 unsatisfactory tests represent a 19% rejection rate based on present preliminary quality control parameters used for the PT-virus.

Out of the 3618 test compounds, 310 demonstrated antiviral activity at greater than 50% reduction levels. This represents around 9% of the tested compounds having in vitro antiviral activity against PTvirus. The remainder, 3308 compounds (91%), are to be considered inactive with present quality control and assay protocols.



Represents 14-month period (November 15, 1989 - January 31, 1991) Figure 62

310

3308

- 4.2.3.1 Prescreen PT-Quality Controls: Two positive control compounds (Ribavirin and 2-Thio-6 Azauridine) were used in the daily assay sets as antiviral activity quality controls. The antiviral performance of the unknown compounds is compared to that of the positive control compounds. Compounds with equal to or better antiviral potency are considered active and are worthy of further in vitro profile studies and in vivo testing.
- 4.2.3.1.1 Antiviral Activity of Ribavirin vs PT Virus A summary of the antiviral and cytotoxicity performance of the primary control compound, AVS-0001 (Ribavirin) is presented in Figure 63-A for 142 tests performed during November, 1989 through January, 1991.

Control Compound-Antiviral Performance: Ribavirin (AVS-0001) has been the sole control compound against PT in these MTT-assay prescreens. The mean and median antiviral inhibition and cytotoxicity patterns of the positive control drug (Ribavirin) are illustrated in Figure 63-A.

The 142 control tests performed with Ribavirin gave a mean Selectivity Index of 13.62 (SD \pm 8.00) and the median value was 12.30 (range = 0.39 - 32.00), indicating moderate antiviral selectivity for Ribavirin. SI is calculated by dividing the TC_{25} by the IC_{50} .

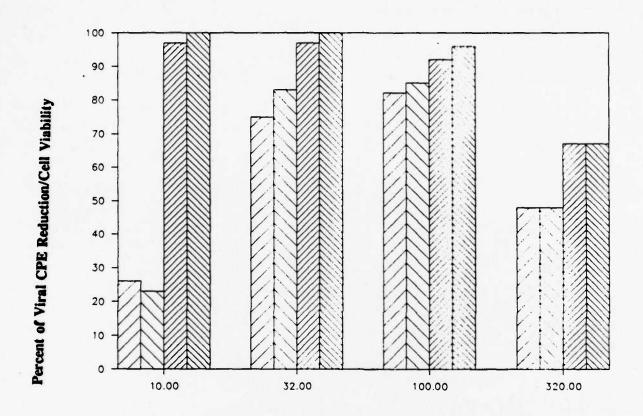
The mean Antiviral Inhibitory Concentration 50% (IC₅₀) was 27.60 μ g/ml (SD \pm 40.90). The median IC₅₀ value was 17.0 μ g/ml (range = 0 - 320). This indicates that Ribavirin does reach 50% antiviral reduction levels, relatively consistent at 10 - 320 μ g/ml (half \log_{10}) concentrations. During this reporting period, the highest and lowest concentration of Ribavirin was varied from 1 to 320 μ g/ml to properly evaluate the maximum antiviral effect and cytotoxicity pattern of Ribavirin. The statistics are based on 142 values tested using the 10 - 320 half \log_{10} scale.

The average maximum antiviral inhibitory level of 142 Ribavirin tests (Figure 63-A) was reached at 100 μ g/ml of the compound with 85% antiviral effect. Maximum antiviral effect (~85%) was found with a simultaneous ~5% cytotoxic suppression. Above this concentration (100 μ g/ml) Ribavirin starts to lose its antiviral potency with increasing cytotoxicity. An increase of the concentration of Ribavirin to 320 μ g/ml, does not improve the antiviral activity (Figure 63-A). The antiviral activity decreases from 85% (at 32 μ g/ml) to 48% (at 320 μ g/ml). The highest concentration (320) of Ribavirin is needed to properly evaluate the Cellular Toxicity 25% (TC₂₅) value.

As reported previously in the 9th semiannual report, a different maximum antiviral value is obtained depending upon which concentration scale is used. In those tests, the \log_{10} scale of 100 - 0.1 measured more accurately the antiviral effect of the control compound, Ribavirin, whereas the \log_{10} scale of 320-0.32 measured more accurately the cytotoxicity effect.

In the present 142 control assays, we tested Ribavirin at 0.5 \log_{10} scale from 10 - 320 μ g/ml concentrations to maximize the correct measurements of both the antiviral and cytotoxicity effects. This enables us to monitor our quality control parameters more accurately.

RIBAVIRIN -VS- PT VIRUS (PRESCREEN PROTOCOL)



CONCENTRATION (µg/ml)

Wean % Viral CPE Reduction	%	Vi Re	edian % ral CPE eduction E Reduction		Cell Vi	ability	Cell Viab	ility	
Conc.(µg/ml)	10	32	100	320	10	32	100	320	
Mean	26	75	82	48	97	97	92	67	
Median	23	83	85	48	100	100	96	67	
Std. Dev.	0.19	0.24	0.15	0.05	0.06	0.10	0.18		

Figure 63-A

Average Antiviral and Cytotoxicity Values for 142 Positive Control Compounds Assays

323

4.2.3.1.2 <u>Maximum Antiviral Effect of Ribavirin vs PT Virus</u>: Since the metabolic activity of the cells was an unknown function during the testing period, it was monitored indirectly by measuring the maximum antiviral effect of the control compound Ribavirin. This demonstrated the amount of infectious virus that was produced by the cells (Maximum Percent CPE).

A bar graph scatter plot (Figure 64-A) depicts the distribution of the maximum antiviral reduction values of all 142 control compound prescreen assays for Ribavirin. The results indicate that the average maximum antiviral reduction obtained with the present SOP is around 88% (SD \pm 12.00) reduction levels. The maximum reduction levels vary from 56 - 100% but remain quite consistently around the median of 90%. The assay control values give a shifted half-bell-shaped distribution curve toward the maximum 100% reduction level. This indicates quite a consistent day-to-day performance of the control compound in the PT prescreen-MTT assay.

During this period the positive control compound performance criteria for Ribavirin versus the PT virus in the pre-screen format has not been set to a definite endpoint. We have collected data in order to find out what would constitute a reasonable quality control endpoint and the data indicates that it could be set at the 50% reduction level. In order to measure the maximum antiviral endpoints of Ribavirin correctly, the concentration scale and the highest concentration to be used, must be evaluated at the proper (semi-log) scale as seen in Figure 63-A.

Ribavirin is active in vitro against PT virus and functions as a reasonable quality control compound. On the other hand, regardless of the performance of the PT-quality control drug Ribavirin, around 32 other compounds have equal or better antiviral activity against PT virus than AVS-0001.

Variation of the Maximum Antiviral Effect PT Virus - VS - Ribavirin (Prescreen Protocol)

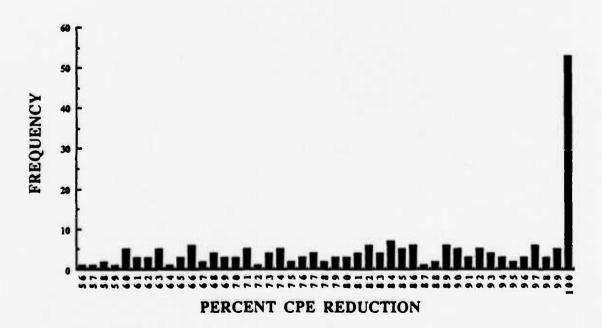


Figure 64-A

Maximum Antiviral CPE Reduction (%).

Summary of 142 Control Tests.

4.2.3.1.3 Cellular Cytotoxicity of Ribavirin vs PT Virus

<u>PT-Control Compound-Cytotoxicity Performance</u>: The 142 cytotoxicity values of the positive control compound Ribavirin are also very consistent. The mean cell **Toxic Concentration 25%** (TC₂₅) was 240 μ g/ml (SD \pm 81.60) and the median was 262 μ g/ml (range of 8.62 to 320 μ g/ml). The reason for this discrepancy is that at 320 μ g/ml scale, the TC₂₅ cytotoxicity cannot always be measured accurately.

As can be seen from Figure 63-A, a definite TC_{25} toxicity value can be measured with a 320 μ g/ml log_{10} scale. Further increase in the concentration of Ribavirin would be needed to consistently evaluate the maximum cytotoxicity of Ribavirin.

Also Figure 63-A, indicates that when the cytotoxicity reaches $\sim 5\%$ at 100 μ g/ml, the control compound (Ribavirin) has reached simultaneously its maximum antiviral effect. The cytotoxic effect of Ribavirin is insignificant between 1 and 100 μ g/ml. The average cytotoxicity reached 37% at 320 μ g/ml, which was the highest Ribavirin concentration tested.

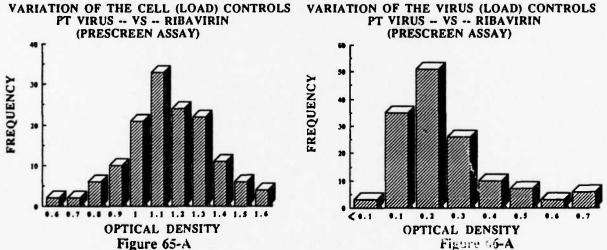
Ribavirin has a definite cytotoxic suppression on cellular metabolism and growth. However, the TC_{25} and TC_{50} toxicity could not be consistently achieved with the 100 μ g/ml concentration of Ribavirin. Therefore, a readjustment to 320 μ g/ml (as being the highest Ribavirin concentration tested) was done during this reporting period. However, at this concentration (320 μ g/ml) the TC_{50} and TC_{95} cannot yet be measured consistently.

4.2.3.1.4 PT-Assay Plate Ouality Controls: Cell Load and Virus Load Parameters (Ribavirin) The MTT assay is fundamentally dependent upon the quality of the assay plates. Our large-scale antiviral testing is dependent upon the uniformity of the test plates produced for the daily assays. Equal numbers of cell load and virus load as well as the consistent performance of the reagents used daily was monitored. A sample of the plate variation control for the period of November, 1989 through January, 1991, is presented in Figures 65-A, 66-A and 67-A.

<u>PT-Control Compound-Cell Load Performance</u>: A bar graph scatter plot of the mean cell control (O.D. reading) of 142 control assays is plotted in Figure 65-A. The results indicate that the cell O.D. readings reached a mean $1.150 \text{ (SD} \pm 0.200)$ with a median of 1.130 (range of 0.610 - 1.630). This indicates that a uniform and equal number (18,000 cells/well) of cells are being loaded into every well in the 96-well plate during the day-to-day operation. The cells reduced MTT to formazan giving maximum blue color uniformly and consistently.

<u>PT-Control Compound-Virus Load Performance</u>: A bar graph scatter plot of the mean virus load O.D. readings of the 142 control assays is presented in Figure 66-A. The results indicate that the average virus load O.D. reading is 0.240 (SD \pm 0.150) with a median of 0.200 (range of 0.020 - 0.730). This demonstrates that a reasonable cell destruction is taking place and a uniform load of virus (32 TCID₅₀) is administered on the cell monolayer with consistent viral CPE results.

PT-Control Compound-Assay Differential Performance: A bar graph scatter plot of the mean O.D. differential values of the 142 control assays is provided in Figure 67-A. The results indicate that the average differential O.D. reading is 0.897 (SD \pm 0.211) with a median of 0.214 (range 0.460 - 1.423). The single bell-shaped curve is reasonably sharp and uniform. This reflects that the assays are executed consistently and are repeatable during day-to-day operation with close to 89% measurement accuracy.



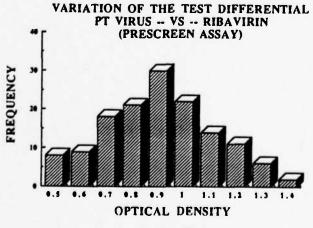


Figure 67-A

4.2.3.1 Prescreen PT-Ouality Controls:

4.2.3.1.1 Antiviral Activity of 2-Thio-6-Azauridine vs PT Virus: A summary of the antiviral and cytotoxicity performance of the second control compound, AVS-6724 (2-Thio-6-Azauridine) is presented in Figure 63-B for 60 tests performed during November, 1989 through January, 1991.

Second Control Compound-Antiviral Performance: 2-Thio-6-Azauridine (AVS-6724) has been tested as a possible second control compound against PT in these MTT-assay prescreens. The mean and median antiviral inhibition and cytotoxicity patterns of this second positive control drug are illustrated in Figure 63-B.

The 60 control tests performed with 2-Thio-6-Azauridine gave a mean Selectivity Index (SI) of 17.40 (SD \pm 8.20) and the median value was 18.00 (range = 0.30 - 32.00), indicating moderate antiviral selectivity for 2-Thio-6-Azauridine. The reason for this discrepancy is that at 100 μ g/ml the 25% cytotoxicity cannot be measured properly to execute SI calculations. SI is calculated by dividing the TC₂₅ by the IC₅₀.

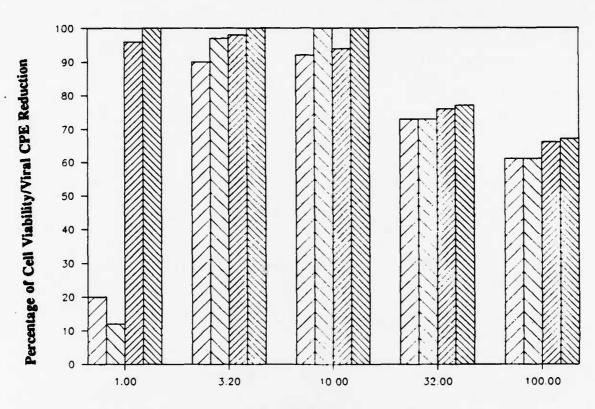
The mean Antiviral Inhibitory Concentration 50% (IC₅₀) was 2.00 μ g/ml (SD \pm 0.92). The median IC₅₀ value was 1.7 μ g/ml (range = 1 - 5.35). This indicates that 2-Thio-6-Azauridine does reach 50% antiviral reduction levels consistently at 32 μ g/ml maximum concentration. During this reporting period, the highest concentration of 2-Thio-6-Azauridine was varied from 32 to 320 μ g/ml to properly evaluate the maximum antiviral effect and cytotoxicity pattern of 2-Thio-6-Azauridine.

The average maximum antiviral inhibitory level of 60 2-Thio-6-Azauridine tests (Figure 63-B) was reached at 3.2 μ g/ml of the compound with 95% antiviral effect. Maximum antiviral effect (~95%)was found with a simultaneous ~2% cytotoxic suppression. Above this concentration (10 μ g/ml) 2-Thio-6-Azauridine starts to lose its antiviral potency with increasing cytotoxicity (Figure 63-B). An increase of the concentration of 2-Thio-6-Azauridine to 100 μ g/ml, does not improve the antiviral activity (Figure 63-B). The antiviral activity decreases from 92% (at 10 μ g/ml) to 60% (at 100 μ g/ml).

A different maximum antiviral value is obtained depending upon which concentration scale is used. In these tests (Figure 63-B), the \log_{10} scale of 1 - 100 measures more accurately the antiviral effect of the control compound, 2-Thio-6-Azauridine, whereas the \log_{10} scale of 0.32 - 320 μ g/ml measures more accurately the cytotoxicity effect.

In the present 60 positive control assays, we tested 2-Thio-6-Azauridine at 0.5 \log_{10} scale concentrations (1 - 32 μ g/ml) to maximize the correct measurements of both the antiviral and cytotoxicity effects. This enables us to monitor our quality control parameters more accurately.

2-THIO-6-AZAURIDINE -VS- PT (PRE-SCREEN) VIRUS



CONCENTRATION (µg/ml)

Wiral CPE
Reduction

Median % Viral CPE Reduction Cell Viability

Median %
Cell Viability

% Viral CPE Reduction

% Cell Viability

Conc.(µg/ml)	\$6. P	1	3.2	10	32	100	1	3.2	10	32	100
Mean		4	35	79	69	73	98	99	96	79	80
Median		0	36	80	76	72	100	100	100	81	82
Std. Dev.	0.	09	0.24	0.19	0.23	0.08	0.04	0.03	0.06	0.14	0.08

Figure 63-B
Average Antiviral and Cytotoxicity Values for 60 Positive Control Compound Tests

4.2.3.1.2 <u>Maximum Antiviral Effect of 2-Thio-6-Azauridine vs PT Virus:</u> Since the metabolic activity of the cells was an unknown function during the testing period, it was monitored indirectly by measuring the maximum antiviral effect of the control compound 2-Thio-6-Azauridine. This demonstrated the amount of infectious virus produced by the cells (Maximum Percent CPE).

A bar graph scatter plot (Figure 64-B) depicts the distribution of the maximum antiviral reduction values of all 60 control compound prescreen assays for 2-Thio-6-Azauridine. The results indicate that the average maximum antiviral reduction obtained with the present SOP is around 97.60% (SD \pm 5.30) reduction levels. The maximum reduction levels vary from 77 - 100% but remain quite consistently around the median of 100%. The assay control values give a shifted half-bell-shaped distribution curve toward the maximum 100% reduction level. This indicates quite a consistent day-to-day performance of the second control compound in the PT prescreen-MTT assay.

During this period the positive control compound performance criteria for 2-Thio-6-Azauridine versus the PT virus in the pre-screen format has not been set to a definite endpoint. We have collected data in order to find out what would constitute a reasonable quality control endpoint and the data indicates that it could be set at the 50% reduction level. In order to measure the maximum antiviral endpoints of 2-Thio-6-Azauridine correctly, the concentration scale and the highest concentration to be used, must be evaluated at the proper semi-log scale as seen in Figure 63-B.

2-Thio-6-Azauridine is active *in vitro* against PT virus and functions as a reasonable candidate for a second quality control compound.

Variation of the Maximum Antiviral Effect PT Virus - VS - 2-Thio-6-Azauridine (Prescreen Protocol)

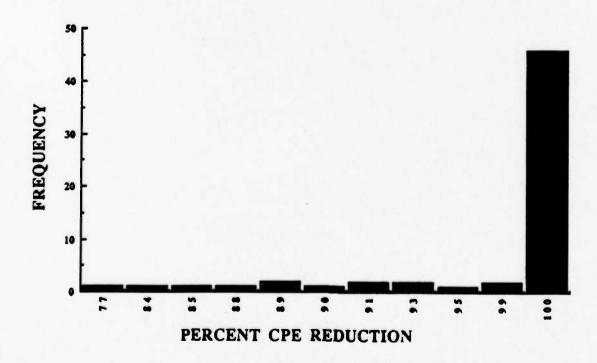


Figure 64-B

Maximum Antiviral CPE Reduction (%).

Summary of 60 Control Tests.

4.2.3.1.3 Cellular Cytotoxicity of 2-Thio-6-Azauridine vs PT Virus

<u>PT-Control Compound-Cytotoxicity Performance</u>: The 60 cytotoxicity values of the positive control compound 2-Thio-6-Azauridine are also very consistent. The mean cell **Toxic Concentration** 25% (TC₂₅) was 32.9 μ g/ml (SD \pm 20.70) and the median was 32.00 μ g/ml (range of 0.88 to 100 μ g/ml). The reason for this discrepancy is that at 100 μ g/ml scale the TC₂₅ cytotoxicity cannot be measured accurately.

As can be seen from Figure 63-B, a definite TC_{25} toxicity value can be measured with a 100 μ g/ml 0.5 log_{10} scale. Further increase in the concentration of 2-Thio-6-Azauridine would be needed to properly evaluate the maximum cytotoxicity of 2-Thio-6-Azauridine.

Also Figure 63-B, indicates that when the cytotoxicity reaches ~ 0 - 5% at 10 μ g/ml, the control compound (2-Thio-6-Azauridine) has reached simultaneously its maximum antiviral effect. The cytotoxic effect of 2-Thio-6-Azauridine is insignificant between 0 and 10 μ g/ml. The average cytotoxicity reached 35% at 100 μ g/ml, which was the highest 2-Thio-6-Azauridine concentration in most tests.

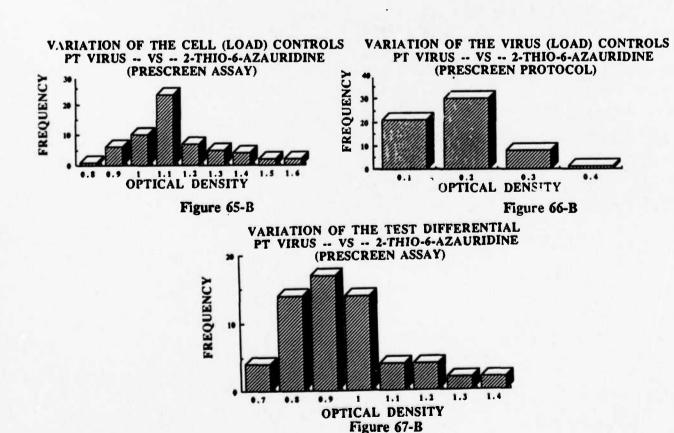
2-Thio-6-Azauridine has a definite cytotoxic suppression on cellular metabolism and growth. However, the TC_{25} toxicity could not be consistently achieved with the 32 μ g/ml concentration of 2-Thio-6-Azauridine. Therefore, a readjustment to 100 μ g/ml (as being the highest 2-Thio-6-Azauridine concentration tested) would be needed to properly evaluate the TC_{25} endpoint. However, at this concentration (100 μ g/ml) the IC_{50} cannot be measured.

4.2.3.1.4 PT-Assay Plate Quality Controls: Cell Load and Virus Load Parameters (2-Thio-6-Azauridine): The MTT assay is fundamentally dependent upon the quality of the assay plates. Our large-scale antiviral testing is dependent upon the uniformity of the test plates produced for the daily assays. Equal loads of cell load and virus load as well as the consistent performance of the reagents used daily was monitored. A sample of the plate variation control for the period of November, 1989 through January, 1991 is presented in Figures 65-B, 66-B, and 67-B.

<u>PT-Control Compound-Cell Load Performance</u>: A bar graph scatter plot of the mean cell control (O.D. reading) of 60 control assays is plotted in Figure 65-B. The results indicate that the cell O.D. readings reached a mean 1.140 (SD \pm 0.170) with a median of 1.100 (range of 0.840 - 1.640). This indicates that a uniform and equal number (18,000 cells/well) of cells are being loaded into every well in the 96-well plate during the day-to-day operation. The cells reduced MTT to formazan giving maximum blue color uniformly and consistently.

<u>PT-Control Compound-Virus Load Performance</u>: A bar graph scatter plot of the mean virus load O.D. readings of the 60 control assays is presented in Figure 66-B. The results indicate that the average virus load O.D. reading is 0.190 (SD \pm 0.070) with a median of 0.160 (range of 0.100 - 0.450). This demonstrates that a reasonable cell destruction is taking place and a uniform load of virus (32 TCID₅₀) is administered on the cell monolayer with consistent viral CPE results.

PT-Control Compound-Assay Differential Performance: A bar graph scatter plot of the mean O.D. differential values of the 60 control assays is provided in Figure 67-B. The results indicate that the average differential O.D. reading is 0.950 (SD \pm 0.170) with a median of 0.940 (range 0.687 - 1.423). The single bell-shaped curve is reasonably sharp and uniform. This reflects that the assays are executed consistently and are repeatable during day-to-day operation with close to 94% measurement accuracy.



4.2.3.2 Prescreen PT-Antiviral Activity Results:

New Drugs with 50% Antiviral Reduction Levels: Out of the 3618 accepted single drug tests, 310 new compounds demonstrated antiviral activity, having antiviral reduction values equal to or better than 50%. This represents around 8.6% of the test compounds being active at this antiviral reduction level. These compounds are summarized in Table 33 according to the highest Selectivity Index (SI). Four compounds, B721392, B721377, B721595 and B723280 demonstrated the best antiviral promise having SI's of > 1000. Seventeen other compounds demonstrated good antiviral activity, having SI's that ranged from 102-883. The rest (289 compounds) showed marginal to moderate antiviral activity with SI's that ranged from 0.2 to 88.

Table 33

New Prescreen Drugs that Produced 50% Antiviral Reduction Against PT Virus

-							
DRUG		PLT	SHIP	TEST			
#	VIR	#	#	DATE	TC 25	1C 50	SI
B721392	PT	V5C	10	0/ /12 /00	- 4000 O	4 00 5 1	000 0
		V59		04/12/90			0.00
8721377	PT			04/12/90			000.0
B72 1595	PT	V63	4P	04/13/90			0.00
8723280	PT	XP4		07/24/90			000.0
8723278	PT	XP3		07/24/90		< 1.00 > 8	
8721374	PT	V59		04/12/90		< 1.00 > 5	
8722904	PT	VOM		04/26/90		< 1.00 > 5	
8721611	PT		4P	04/13/90	408.00		07.65
8721596	PT	V63		04/13/90		< 1.00 > 3	
8721011	PT	VU9		05/01/90	328.00		27.84
8720952	PT	U25	4P	04/03/90	326.00	< 1.00 > 3	
8720951	PT	UZ5	4P	04/03/90	326.00	< 1.00 > 3	25.56
8722805	PT	UBK		03/07/90	325.00	< 1.00 > 3	
8721378	PT	V59	4P	04/12/90	318.00	< 1.00 > 3	18.26
8721021	PT	VUB	4P	05/01/90	265.00	< 1.00 > 2	64.97
8720958	PT	UZ6	4P	04/03/90	231.00	< 1.00 > 2	31.38
8722874	PT	VH6	3P	04/17/90	701.00	3.20 2	19.04
8720916	PT	UZ4	4P	04/03/90	170.00	< 1.00 > 10	69.54
B720937	PT	UZ4	4P	04/03/90	169.00	< 1.00 > 1	68.79
8720963	PT	U26	4P	04/03/90	168.00	< 1.00 > 1	68.28
8720941	PT	U25	4P	04/03/90	102.00	< 1.00 > 1	02.16
8722871	PT	VH6	3P	04/17/90	334.00	3.78	88.45
8721568	PT	09N		08/03/89	330.00	5.77	57.24
8722808	PT	UBK	3P	03/07/90	35.10	< 1.00 > 3	35.14
8721568	PT	089		07/19/89	290.00	8.50	34.08
8724825	PT	0EM		09/14/89	156.00	4.75	32.91
8724374	PT	06H		06/29/89	1000.0		31.60
B849179	PT	X10	6P	07/13/90	701.00		29.68
8724526	PT	095		07/26/89	161.00	5.60	28.78
8724447	PT	07K		07/12/89	208.00		28.58
8722054	PT	OIW		11/08/89	786.00		27.94
8720940	PT	U25	4P	04/03/90	27.80	< 1.00 >	27.76
8720979	PT	UZ6	4P	04/03/90			27.59
8722077	PT	01X		11/08/89	211.00	7.88	26.75
8721064	PT	VRA	4P	04/27/90	742.00	34.80	21.33
8724740	PT	OCY		08/30/89	76.60	3.78	20.27
8723037	PT	UHH	3P	03/14/90	> 1000.0	51.10 >	19.56
8721062	PT	VR9	4P	04/27/90	595.00	32.90	18.10
8721749	PT	06C		06/29/89	1000.0	59.40	16.80
8721702	PT	05V		06/21/89	1000.0	61.00	16.40
8721061	PT	VR9	4P	04/27/90	409.00	29.10	14.09
8722228	PT	OK1		11/22/89	487.00		14.07
8722081	PT	OIY		11/08/89	134.00	9.54	14.03
8721880	PT	OGV		10/11/89	> 1000.0	71.50 >	13.99
8721568	PT	0A4		08/09/89	241.00	17.80	13.49
8722222	PT	OKO		11/22/89	446.00	34.30	13.02
8721092	PT	09K		08/03/89	325.00		12.35
8721511	PT	09N		08/03/89	330.00		11.61
8724521	PT	094		07/26/89	32.80		11.38
8724698	PT	OCF		08/24/89	51.50		11.19
8722091	PT	012		11/08/89	310.00		11.17
8721592	PT	V63	4P	04/13/90	330.00		11.00
8721690	PT	05T		06/21/89	1000.0		10.60
8722525	PT	TE7	3P	01/18/90	323.00		10.53

Table 33 (Cont'd)

### REPAIRS PT OPM	DRUG	VIR	-	SHIP	TEST DATE	TC 25	IC 50	SI
4279 PT 0994 08/03/89 32.50 3.16 10.28 8724566 PT 0994 08/03/89 32.50 3.36 10.28 8724566 PT 0994 08/03/89 32.50 3.36 0.70 8721756 PT 066 06/29/89 339.00 38.60 8.77 8721756 PT 066 06/29/89 339.00 38.60 8.77 8721756 PT 066 06/29/89 339.00 38.60 8.70 8721754 PT 067 08/24/90 538.00 61.80 8.70 8721754 PT 068 06/29/89 337.00 41.50 8.13 8724688 PT 079 07/12/89 28.90 40.10 8.13 8724688 PT 079 07/12/89 28.90 40.10 8.13 8724688 PT 079 07/12/89 28.90 40.10 7.36 8724588 PT 079 09/27/89 505.00 64.40 7.36 8724588 PT 079 09/27/89 505.00 64.40 7.36 8724588 PT 079 09/27/89 505.00 64.40 7.36 8724589 PT 080 39 03/07/90 543.00 71.20 7.63 8722843 PT VBL 39 04/26/90 366.00 49.20 7.43 8724813 PT 081 01/26/90 366.00 49.20 7.43 8724813 PT 081 01/26/90 366.00 49.20 7.43 8724813 PT 081 01/26/90 306.00 67.50 7.52 8724920 PT 251 10 10/26/90 306.00 44.50 7.31 8724813 PT 081 01/25/89 1000.0 137.00 > 7.29 8724416 PT 068 07/06/89 29.40 4.09 7.17 8724813 PT 081 01/25/89 1000.0 137.00 > 7.29 8724416 PT 068 07/06/89 29.40 4.09 7.17 88349035 PT XIH 69 07/13/90 115.00 16.20 7.10 8724820 PT 094 07/26/89 29.40 4.09 7.17 88349035 PT XIH 69 07/13/90 150 00 63.70 6.83 8724433 PT 074 07/26/89 22.70 3.65 60 4.21 6.79 8724420 PT 09L 09/14/89 23.40 3.61 6.48 872453 PT 074 07/26/89 23.60 3.61 6.40 7.70 8724769 PT 0824799 49.00 7.64 6.0 6.47 8722476 PT 0824799 49.00 7.64 6.0 6.27 8722476 PT 0824799 49.00 7.64 6.0 6.37 6.0 5.38 8722473 PT 091 018 11/08/89 325.00 55.60 5.80 5.50 5.80 8722478 PT 091 018 11/08/89 325.00 55.60 5.80 5.50 5.80 8722478 PT 092 07/26/89 303.00 6.61 6.0 5.28 872478 PT 092 07/26/89 303.00 6.61 6.0 5.28 872478 PT 093 000 000 11/26/89 500 000 5.59 5.50 5.50 5.80 8722279 PT 001 01/06/89	R774454	DT	0714		07/12/89	74 00	* **	10 48
### R724566 PT								10.28
### ### ### ### ### ### ### ### ### ##	4280							
### PT 06F 06/29/89 339.00 38.60 8.70 ### PT 06F 06/29/89 339.00 37.30 8.70 8.70 ### PT 06F 06/29/89 335.00 37.30 8.70 ### PT 06F 06/29/89 335.00 34.60 4.07 8.50 ### PT 06F 06/29/89 337.00 41.50 8.13 ### PT 06F 06/29/89 337.00 41.50 8.13 ### PT 06F 07/12/89 337.00 41.50 8.13 ### PT 07F 07/12/89 337.00 41.50 8.13 ### PT 07F 07/12/89 337.00 41.50 8.13 ### PT 07F 07/12/89 326.00 40.10 8.13 ### PT 07F 07/26/89 326.00 40.10 7.55 ### PT 07F 07/26/89 326.00 40.10 7.55 ### PT 07F 07/26/89 326.00 46.40 7.84 ### PT 07F 07/26/89 326.00 47.50 7.53 ### PT 07F 07/26/89 326.00 47.50 7.53 ### PT 07F								
### ### ### ### ### ### ### ### ### ##				4P	to the same of the			
B722280 PT Y YGG 7P 08/24/90 38.00 61.80 8.70 B721754 PT YGG 7P 08/24/98 337.00 41.50 8.50 B721754 PT OKE 06/29/89 337.00 41.50 8.13 B7226458 PT OW 11/08/89 332.00 41.50 8.13 B724686 PT OFE 09/27/89 505.00 64.60 7.84 B722847 PT USO 09/27/89 505.00 64.60 7.84 B722843 PT USO 09/27/89 505.00 64.50 7.34 B722843 PT USO 09/26/90 588.00 67.50 7.52 B722843 PT USO 09/26/90 588.00 67.50 7.52 B722823 PT SET 10 10/26/90 586.00 67.50 7.41 GRP19386 PT VSA 4P 04/26/90 330.00 64.50 7.43 B724416 PT OKR 07/06/89 29.40 4.09 7.13 B724416 PT OKR 07/06/								
8722634 PT V06 7P 06/26/90 34.60 4.07 8.50 8721734 PT 06E 06/29/89 337.00 41.50 8.13 8722048 PT 01V 11/08/89 326.00 40.10 8.13 8724586 PT 07B 07/12/89 28.90 3.69 7.24 8724527 PT 066 07/26/89 32.50 4.19 7.75 8722849 PT 100 30 30/70/90 53.00 71.20 7.63 8722883 PT VQL 3P 04/26/90 508.00 67.50 7.52 8721917 PT 0NA 10/18/89 330.00 44.50 7.41 8721823 PT 0NA 10/18/89 330.00 44.50 7.43 8721823 PT 0NS 10/25/89 > 1000.0 137.00 > 7.29 8724164 PT 0KS 07/06/89 29.40 4.09 7.17 872183 PT 0XX 11/22/89 271.00 36.00 67.50 7.3 8724616 PT 0KS 07/06/89 32.00 44.50 7.17 872183 PT 0XX 11/22/89 271.00 36.00 64.50 7.3 8724533 PT 0XX 11/22/89 271.00 36.00 67.73 8724635 PT XIII 6P 07/13/90 115.00 6.37 0.63 8724620 PT 0EL 09/14/89 23.40 3.61 6.83 8724820 PT 0F 0VQ 09/04/89 22.70 3.55 6.58 8724728 PT 0VQ 09/04/89 22.70 3.55 6.58 8724728 PT 0VQ 09/04/89 32.00 50.00 50.00 50.00 6.41 8724735 PT 000				7P				
### 8724458 PT 01V 01V 11/08/89 337.00 41.50 8.13 8724458 PT 01V 01V 11/08/89 336.00 40.10 8.13 8724458 PT 07M 07/12/89 28.90 3.69 40.10 8.13 8724458 PT 07M 07/12/89 28.90 3.69 40.10 8.13 8724527 PT 096 07/26/89 32.50 4.19 7.75 8722849 PT U80 3P 03/07/90 508.00 64.60 7.84 8724527 PT 096 07/26/89 32.50 4.19 7.75 8722849 PT U80 3P 04/26/90 508.00 67.50 7.52 8722920 PT 25T 10 10/26/90 508.00 67.50 7.52 8722920 PT 25T 10 10/26/90 366.00 49.20 7.43 8721917 PT 0NA 10/18/89 330.00 44.50 7.41 6RP19386 PT V8X 4P 04/26/90 326.00 44.50 7.41 6RP19386 PT 0NS 10/25/89 20.40 44.50 7.35 8721823 PT 0NS 10/25/89 271.00 38.00 7.35 8721823 PT 0NS 11/22/89 271.00 38.00 7.35 872435 PT 0NS 11/22/89 271.00 38.00 7.37 872183 PT 0NS 11/22/89 271.00 38.00 7.37 8724519 PT 0NA 07/12/89 28.60 4.21 6.79 8724519 PT 0NA 07/12/89 28.60 4.21 6.79 8724520 PT 0EL 09/14/89 23.40 3.61 6.85 8724520 PT 0EL 09/14/89 23.40 3.61 6.86 8722632 PT VB 7P 08/24/90 494.00 76.40 6.47 8724728 PT 052 06/21/89 1000.0 156.00 6.41 8724728 PT 052 06/21/89 350.00 3.74 6.41 8848752 PT 21S 11P 01/17/91 3200.0 501.00 > 6.38 8724453 PT 000 09/06/89 24.00 3.74 6.41 8848752 PT 21S 11P 01/17/91 3200.0 501.00 > 6.38 8724453 PT 000 09/06/89 24.00 3.64 6.41 8724578 PT 000 08/30/89 22.30 3.64 6.12 8724466 PT 070 07/12/89 355.00 55.60 5.84 8724477 PT 06R 07/06/29 25.10 3.80 5.81 8721758 PT 06L 08/30/89 325.00 55.60 5.54 6.72 8724733 PT 00H 09/06/89 21.50 3.86 5.50 5.55 8724783 PT 06L 08/30/89 325.00 55.60 5.54 6.72 872473 PT 06L 08/30/89 325.00 55.60 5.54 6.72 872473 PT 06G 08/29/89 300.0 0 185.00 5.39 8721761 PT 06G 08/29/89 300.0 0 55.00 55.50 5.55 8724584 PT 06C 08/29/89 300.0 0 55.00 55.50 5.55 8724584 PT 06C 08/29/89 300.0 0 55.00 55.50 5.55 8724584 PT 06C 08/29/89 300.0 0 55.00 55.50 5.55 8724584 PT 06C 08/29/89 300.0 0 55.00		-	-					
B724458 PT OPN O7/12/89 28.90 3.69 7.84 B724527 PT OPS 09/27/89 505.00 64.40 7.86 B722527 PT OPS 07/26/89 32.50 4.19 7.85 B722883 PT VOL 3P 03/07/90 508.00 67.50 7.52 B722920 PT 2ST 10 10/26/90 366.00 49.20 7.35 B722920 PT 2ST 10 10/26/90 366.00 49.20 7.41 GRP19386 PT VPX 4P 04/26/90 366.00 44.50 7.36 B724416 PT OGR 07/06/89 29.40 4.09 7.13 B722833 PT VIII 6P 07/06/89 29.40 4.09 7.13 B722886 PT VIII 3P 0/16/89 22.40 4.09 7.13 B722519 PT OUR 3P 0/12/89 22.10	B721754	PT	06E					8.13
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B724762 PT OOC 09/06/89 20.40 4.13 4.95 B724712 PT OCH 08/24/89 19.80 4.14 4.77 8848750 PT IEE 11P 11/29/90 > 1000.0 212.00 > 4.72 8724434 PT 07H 07/12/89 30.30 6.61 4.58 8724586 PT 0A6 08/09/89 208.00 45.60 4.57 8724457 PT 07M 07/12/89 16.20 3.57 4.55 8724387 PT 090 07/26/89 263.00 57.80 4.55 8724384 PT 06K 07/06/89 23.40 5.21 4.49 8724418 PT 06R 07/06/89 18.80 4.24 4.44 87224670 PT 0C9 08/24/89 66.40 14.90 4.44 8722080 PT 0K1 11/22/89 192.00 47.30 4.07 8721719 PT </td <td></td> <td></td> <td></td> <td></td> <td></td> <td>-</td> <td></td> <td></td>						-		
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8848750 PT 1EE 11P 11/29/90 1000.0 212.00 4.72 B724434 PT 07H 07/12/89 30.30 6.61 4.58 B724586 PT 0A6 08/09/89 208.00 45.60 4.57 B724457 PT 07M 07/12/89 16.20 3.57 4.55 B724387 PT 090 07/26/89 263.00 57.80 4.55 B724384 PT 06K 07/06/89 23.40 5.21 4.49 B724418 PT 06R 07/06/89 18.80 4.24 4.44 B724670 PT 0C9 08/24/89 66.40 14.90 4.44 B722230 PT 0K1 11/22/89 192.00 47.30 4.07 B722080 PT 0IY 11/08/89 108.00 26.70 4.05 B721719 PT 05X 06/21/89 100.00 266.00 3.76 B724558 PT </td <td></td> <td></td> <td></td> <td></td> <td>9.00</td> <td></td> <td></td> <td></td>					9.00			
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B724670 PT OC9 08/24/89 66.40 14.90 4.44 B722230 PT 0K1 11/22/89 192.00 47.30 4.07 B722080 PT 0IY 11/08/89 108.00 26.70 4.05 B721719 PT 05X 06/21/89 1000.0 266.00 3.76 B724607 PT 0AB 08/09/89 18.80 5.01 3.76 B724558 PT 090 08/03/89 26.10 6.97 3.74 B721838 PT 0GS 10/11/89 212.00 56.80 3.73 B724508 PT 087 07/19/89 20.00 5.49 3.64 B722889 PT VQL 3P 04/26/90 348.00 97.30 3.57								
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B722889 PT VQL 3P 04/26/90 348.00 97.30 3.57	The state of the s							
	B722889			3P			The second second	
	B721910	PT	0H9			1000.0	281.00 >	

Table 33 (Cont'd)

ORUG #	VIR		SHIP	TEST OATE	TC 25	IC 50	SI
		0/1				202.00	
8721734 8724423	PT PT	06A 06S		06/29/89 07/06/89	1000.0 12.20	282.00 3.47	3.55 3.53
8724382	PT	061		07/06/89	168.00	47.80	3.51
8724413	PT	060		07/06/89	15.50	4.43	3.49
8722078 8724596	PT	OA9		11/08/89 08/09/89	130.00 324.00	37.30 94.30	3.49
B848669	PT		11P	11/21/90 >		291.00 >	3.44
8724728	PT	OCV		08/30/89	15.20	4.51	3.38
B721979	PT	OHX		10/25/89 >	1000.0	298.00 >	3.35
8724592 8722628	PT	OA8	7p	08/09/89 08/24/90 >	226.00 1000.0	67.70 300.00 >	3.34
8722811	PT	UBL		03/07/90	325.00	98.20	3.31
8721908	PT	049		10/18/89	328.00	99.80	3.29
8724652 8709763	PT PT	087 2CB	13P	08/16/89	199.00	60.70	3.28
B724855	PT	OF4	ISP	01/03/91 > 09/27/89	1000.0 15.90	316.00 > 5.08	3.16 3.13
B724860	PT	OF5		09/27/89	204.00	65.40	3.12
B722241	PT	OK2		11/22/89 >	1000.0	320.00 >	3.12
8722052 8722117	PT	OIV 2DX	00	11/08/89 10/10/90	164.00 218.00	53.00 71.20	3.10 3.07
B724732	PT	OCW	77	08/30/89	29.80	9.73	3.06
B710043	PT	2CD	13P	01/04/91 >	1000.0	329.00 >	3.04
B724719	PT	OCT		08/30/89	14.80	4.98	2.97
8724714 8724549	PT PT	0CS		08/30/89 08/03/89	19.20 285.00	6.61 99.00	2.90
8724553	PT	09P		08/03/89	113.00	39.70	2.84
3593	PT	05W		06/21/89	1.23	0.44	2.82
B724644	PT	085		08/16/89	182.00	65.60	2.77
8724455 8721722	PT PT	07L 05Y		07/12/89 06/21/89	9.95 1000.0	3.61 365.00	2.75
8710057	PT	210	13P	01/17/91	2610.0	962.00	2.72
8724590	PT	OA7		08/09/89	162.00	60.00	2.71
8710110	PT	207	13P	01/08/91 >	1000.0	370.00 >	2.70
B721708 B724863	PT	05W		06/21/89 09/27/89	1000.0 150.00	371.00 55.60	2.69
8721953	PT	OIR		11/08/89 >	1000.0	372.00 >	2.69
8722182	PT	OJX	lucky.	11/22/89	188.00	70.20	2.68
8710128 	PT	208	13P	01/08/91 >	1000.0	374.00 >	2.67
8724724 8724420	PT	0CU 06S		08/30/89 07/06/89	211.00 61.90	82.70 24.50	2.56
8721604	PT	YPK	8P	09/14/90 >	1000.0	399.00 >	2.51
8721755	PT	06E	-	06/29/89	1000.0	400.00	2.50
8848653 8722239	PT	21M OK2	11P	01/16/91 >		1320.0 >	2.42
B722116	PT	ZDW	9P	11/22/89 10/09/90	160.00 196.00	66.50 81.70	2.40
8721731	PT	05Z		06/21/89	1000.0	423.00	2.36
B724664	PT	089		08/16/89 >		425.00 >	2.36
8724394 8724633	PT	06M 083		07/06/89 08/16/89	15.90 185.00	6.80 79.90	2.34
8722246	PT	OK3		11/22/89	171.00	75.80	2.26
8721511	PT	089		07/19/89 >	320.00	144.00 >	2.23
8721714	PT	05X		06/21/89	1000.0	450.00	2.22
8721095 8724785	PT	OOH		08/03/89	184.00 14.80	83.50 6.69	2.21
8724415	PT	06R		07/06/89	8.05	3.66	2.20
8721749	PT	880		07/19/89	120.00	55.10	2.18
8721925 8723297	PT	OHC	40	10/18/89 >	1000.0	463.00 >	2.16
8721680	PT	Y38 055	G	08/21/90 > 06/21/89	1000.0	465.00 > 467.00	2.15
8724373	PT	089		07/19/89	211.00	99.40	2.13
8722823	PT	VIO		08/30/90 >	1000.0	479.00 >	2.09
8848911 8724797	PT	1P4 00J	12P	12/12/90 09/06/89	88.30 191.00	42.20 92.60	2.09
B724627	PT	082		08/16/89	10.90	5.27	2.06
8724729	PT	OCW		08/30/89	9.44	4.63	2.04
8848747	PT	215	11P	01/17/91 >		1570.0 >	2.04
8724373 8722087	PT PT	06H 01Z		06/29/89 >	493.00	250.00 507.00 >	1.97
B724442	PT	07J		07/12/89 >	1000.0	524.00 >	1.91
8721737	PT	06A		06/29/89	1000.0	526.00	1.90
8721743	PT	068		06/29/89	946.00	498.00	1.90
8724436	PT	07H		07/12/89	154.00	81.00	1.90

Table 33 (Cont'd)

DRUG	VIR		SHIP	TEST DATE	TC 25	1C 50	SI
	***			DAIL	10 23		•
B724439	PT	071		07/12/89	145.00	76.00	1.90
8721725 8724385	PT	05Y 06K		06/21/89 07/06/89	1000.0 16.00	529.00 8.60	1.89
B710130	PT		13P	01/08/91 >		539.00 >	1.85
B724396	PT	091		07/26/89	173.00	93.90	1.84
B721826 B722743	PT PT	OG1	70	10/04/89 > 08/28/90 >		550.00 > 549.00 >	1.82
B721348	PT	YOZ		09/13/90 >		558.00 >	1.79
8722745	PT	YHM		08/28/90 >		567.00 >	1.76
8848804	PT	2PB		01/18/91 >		1820.0 >	1.76
B848752 B721730	PT	1EF 05Z	11P	11/29/90 06/21/89	554.00 159.00	317.00 91.40	1.75 1.74
8848989	PT	YIW	7P	08/31/90 >		579.00 >	1.73
B721770	PT	06G		06/29/89	467.00	271.00	1.72
B722141 B721787	PT	OJI OFX		11/15/89 10/04/89	781.00 764.00	461.00 458.00	1.69 1.67
B722873	PT	ZSM	10	10/04/09		600.00 >	1.67
8722929	PT	ZOU	10	10/24/90 >		606.00 >	1.65
B721958	PT	OHS		10/25/89 >		615.00 >	1.63
B722168 B721295	PT	OJK	RP.	11/15/89 09/13/90 >	255.00 1000.0	157.00 612.00 >	1.63 1.63
B710046	PT	211		01/17/91	3170.0	1960.0	1.62
8724379	PT	06J		07/06/89	896.00	565.00	1.59
8724769 8733480	PT	ODE YHI	70	09/06/89 >		630.00 > 523.00	1.59
B722689 B724885	PT	OFA	17	08/28/90 09/27/89	824.00 8.52	5.44	1.58 1.57
B724667	PT	OFV		10/04/89	1.29	0.82	1.57
B848653	PT		11P	11/21/90 >		635.00 >	1.57
B724657 B721166	PT PT	088 09L		08/16/89	820.00 601.00	541.00 402.00	1.52
B724832	PT	OEN		08/03/89 09/14/89	933.00	622.00	1.50 1.50
B706269	PT		13P	01/03/91 >		673.00 >	1.49
B710138	PT		13P	01/08/91 >		674.00 >	1.48
B724544 B722247	PT PT	099 0K3		07/26/89 11/22/89	8.33 103.00	5.65 70.50	1.47
B723290	PT	Y3A	6P	08/21/90 >		680.00 >	1.47
B724720	PT	OCT		08/30/89	9.29	6.37	1.46
B722165	PT	OJK	20	11/15/89 >	1000.0	683.00 >	1.46
8722936 8724535	PT	ZOV 097	10	10/24/90 07/26/89	941.00	650.00 6.75	1.45
B724661	PT	0B9		08/16/89	785.00	570.00	1.38
B724405	PT	06P		07/96/89	7.05	5.15	1.37
B721818 B722006	PT	0G0 011		10/04/89	738.00 1000.0	553.00 758.00 >	1.33
B722867	PT	VOK	3P	04/26/90 >	1000.0	769.00 >	1.30
8848926	Pī	1P8		12/18/90 >	1000.0	771.00 >	1.30
B724509	PT	087		07/19/89	118.00	91.80	1.28
B722559 B724530	PT PT	YGK 096	/P	08/24/90 > 07/25/89	1000.0	787.00 > 94.80	1.27
8848811	PT	2PC	11P	01/18/91 >		2560.0 >	1.25
B723044	PT	UHW	3P	03/14/90 >		809.00 >	1.24
B724642	PT	085		08/16/89	7.77	6.33	1.23
B722162 B724411	PT PT	090		11/15/89 > 07/06/89	1000.0 121.00	815.00 > 99.70	1.23
B724654	PT	087		08/16/89	7.21	5.93	1.21
B723268	PT	XP1	6P	07/24/90 >	1000.0	824.00 >	1.21
B721532	PT	09K		08/03/89	529.00	445.00	1.19
B722089 B724525	PT PT	01Z 095		11/08/89 07/26/89	11.70 7.23	9.83 6.13	1.19
B723062	PT	XOZ	SP.	06/19/90	942.00	800.00	1.18
B724517	PT	094		07/26/89	8.65	7.40	1.17
B724716 B715141	PT PT	OCS 2WG	13p	08/30/89 01/29/91 >	6.78	5.86 884.00 >	1.16
B724866	PT	OF6	135	09/27/89	7.85	7.03	1.12
9848753	PT	215		01/17/91	2470.0	2210.0	1.12
B718582	PT	2XE		01/30/91 >		910.00 >	1.10
8849180 8848671	PT PT	XIO 2IN		07/13/90 01/16/91 >	628.00 3200.0	582.00 3000.0 >	1.08
B710151	PT	20A		01/08/91 >		942.00 >	1.06
B722060	PT	ZDS	99	10/09/90	852.00	811.00	1.05
B722591	PT	YGM	7P	08/24/90 >		959.00 >	1.04
B721892	PT	OGX		10/11/89 >	1000.0	979.00 >	1.02

Table 33 (Cont'd)

DRUG		PLT	SHIP	TEST			
	VIR	#	#	DATE	TC 25	1C 50	\$1
8710179	PT	20 F	13P	01/09/91	771.00	759.00	1.02
B721693	PT	05T		06/21/89	604.00	616.00	0.98
8723431	PT	ZSV	10	10/26/90	723.00	761.00	0.95
B724504	PT	086		07/19/89	6.34	6.89	0.92
B848986	PT	AIA	7P	08/31/90	560.00	628.00	0.89
B720825	PT		12P	12/20/90	881.00	1000.0	0.88
B724400	PT	060		07/06/89	6.40	7.43	0.86
B721605	PT	YPL		09/14/90	673.00	781.00	0.86
B848882	PT		12P	12/11/90	655.00	765.00	0.86
B848841	PT		11P	01/18/91	1280.0	1520.0	0.84
B723471	PT	ZT6	10	10/31/90	51.30	63.50	0.81
B724511	₽T	092		07/26/89	6.93	8.80	0.79
B720911	₽T	YLS	8P	09/08/90	637.00	805.00	0.79
B724847	₽T	0EQ		09/14/89	615.00	787.00	0.78
B849037	PT	XIN	6P	07/13/90	695.00	906.00	0.77
B724656	PT	088		08/16/89	675.00	884.00	0.76
B724900	PT	OFD		09/27/89	59.80	78.90	0.76
B724547	PT	090		08/03/89	6.21	8.33	0.75
B722079	PT	VIO		11/08/89	609.00	814.00	0.75
B724510	PT	092		07/26/89	574.00	788.00	0.73
B724578	PT	0A4		08/09/89	569.00	798.00	0.71
B848881	PT		12P	12/11/90	603.00	868.00	0.69
B724871	PT	0F8		09/27/89	64.30	96.50	0.67
B724631	PT	OB3		08/16/89	594.00	914.00	0.65
B722065	PT	OIW		11/08/89	6.00	9.36	0.64
B848907	₽T	1P4	12P	12/12/90	556.00	932.00	0.60
B721696	PT	05U		06/21/89	295.00	543.00	0.54
B724867	PT	0F7		09/27/89	4.74	9.11	0.52
B721498	PT	YPI		09/14/90	338.00	644.00	0.52
B722614	PT	YGO	7P	08/24/90	491.00	1000.0	0.49
B848880	PT		12P	12/11/90	378.00	781.00	0.48
B721559	PT	09L		08/03/89	370.00	860.00	0.43
B724390	PT	090		07/26/89	3.87	9.20	0.42
B722227	PT	OK1		11/22/89	323.00	769.00	0.42
B722181	PT	OJL		11/15/89	38.90	96.40	0.40
B721740	PT	06B		06/29/89	94.10	275.00	0.34
B722109	PT	OJF		11/15/89	28.20	93.00	0.30
B724853	PT	0ER		09/14/89	201.00	805.00	0.25
B721541	PT	09L		08/03/89	98.80	508.00	0.19
B722824	PT	USM	3P	03/07/90	53.70	316.00	0.17

The *in vitro* antiviral activity of the compounds in Table 33 was further confirmed in most of the compounds. Verification of the antiviral activity of these prescreen actives was tested using the primary screening (confirmatory) protocol. (See Table 34)

4.2.4 Confirmatory Assays (Compounds Selected from Prescreen Testing):

During this reporting period 767 compounds were received (from prescreen testing) for confirmatory testing. If a compound had an SI of ≥ 1 then it was considered as a candidate for confirmatory primary screen testing. The SI of ≥ 1 was only a preliminary endpoint that was being used and was subject to change as more data was generated. Data from the confirmatory assays are summarized in Table 34. Out of 767 confirmatory tests, 515 compounds (67%) were confirmed active during this reporting period. The criteria for activity is that the confirmatory test has to show $\geq 25\%$ reduction in CPE in one or more of the viruses tested. Failure to confirm the activity in these compounds was probably due to differences during the assay conditions:

- 1) In confirmatory assays the concentration range is adjusted to a more accurate semilog scale to maximize the SI and calculate the TAI and it should indicate more accurately the antiviral potential of the test compound.
- 2) Differences in the "differential" of the two runs can cause the compound to read positive or negative, falsely. The variability in the differential can cause false positives or false negative bias in day-to-day testing calculations, reflecting the variability in the maximum activity of the compound.
- 3) The metabolic rate, age, and passage number, etc., of the cells may cause the above observed variability in test compound activity.
- 4) Problems associated with stability and storage of the compound (i.e., different lot numbers, solubility, light sensitivity, hygroscopic, etc.).
- 5) Problems associated with technical execution of large numbers of plates by different technicians.
- 6) During the beginning of the prescreen testing (Shipment 1P 5P), the assays were performed with confluent and stationary cellular monolayers. This procedure has a tendency to create false positive results as compared to the confirmatory assay results. In confirmatory testing actively metabolizing subconfluent monolayers have been used.
- 7) Two different MTT protocols are being compared. The prescreen protocol was prophylactic when we were testing drug shipments 1P 5P. The drug was delivered to the cells before final addition of 62 TCID_{50} of virus to total volume of $200 \mu l$. The confirmatory protocol is therapeutic. The virus 32 TCID_{50} is delivered onto the cells before the addition of the drug in total volume of $100 \mu l$. In later drug shipments the same protocol was used for both prescreen and confirmatory assays.

The results seem to indicate that prophylactic treatment with confluent stationary monolayers causes inconsistent cell infections and therefore causes numerous drugs to read as false positive. Later prescreen testing (starting with shipments 5P) was done with the same therapeutic protocol in order to evaluate more properly the correlation of actives from the prescreen to those of the confirmatory (primary screen) results. The conflicting results should be retested based upon the availability of the compound.

Recommendations of Prescreen Confirmed-Actives Based Upon the In Vitro Results with MTT Assay (Vero Cells)

Based upon the above mentioned variable conditions in assay procedures, which are difficult to control on a -day-to-day basis, we recommend the following:

- 1) Since the prescreen assay protocol has a tendency to produce false positive results, only compounds that demonstrate at least 50% reduction in viral CPE and a selectivity index of ≥ 1.0 will advance to confirmatory testing.
- 2) In order not to miss a possible active lead from the prescreen assay, some nontoxic compounds with an SI of ≤ 1.0 need to be retested.
- 3) Compounds with broad-range activity close to 50% reduction must also be retested in order not to miss an active compound that is not clearly dose responsive.
- 4) In confirmatory assays, compounds having antiviral activity at the "3" activity grade level or consistent repeatable grade "2" level are considered active candidate compounds.

Table 34

Confirmatory Testing of Compounds Selected From Prescreen Testing

0 NT 2
0 0 7
1 0
0
70077
59
4 0
8
B631963-F015

Results of Prescreen Testing*

																-					
W	0	0	0	0	0	0	0	0	0	0	TN	0	NT	N	F	F	¥	N.	N	N	N
SF	2	0	0	0	0	0	1	1	0	0	1	2,0	2	1	0	0	0	1	0	0	0
Ä	0	0	0	0	0	0	0	0	0	0	2	1,0	0	0	0	1	0	0	0	0	0
VE	0	0	0	1	0	0	1	1	1	1	1	0,0	2	0	0	0	0	1	0	1	1
æ	0	0	0	0	0	0	0	0	0	0	0	0,0	0	0	0	0	0	0.	0	0	0
YF	1	0	2	0	0	0	1	2	0	1	1	0,1	0	0	0	0	0	0	0	0	0
AVS No.	9121	9122	9123	9124	9125	9126	9127	9128	9129	9130	11047	7323	11048	11049	11050	11051	11052	11053	11054	11055	11056
Z	25	26	11	1	9	14	18	16	25	26	52	33	32	15	29	52	28	35	24	22	23
VE	1	3	4	10	111	22	3	3	4	21	14	1	02	2	2	13	0	1	3	12	12
YF	1	0	40	0	0	1	1	0	0	7	15	14	0	21	1	10	0	2	0	3	1
Prescreen No.	B714994	B714997	B715001	B715010	B715011	B715012	B715022	B715023	B715026	B715060	B715141	B716493-D021	B718574	B718577	B718580	B718582	B718583	B718586	B718587	B718595	B718598
Shipment No.	12P	13P	4P	14P																	

Results of Prescreen Testinga

Prescreen YF VE FT AVS YF JE VE FT S										- 11		
B718599 0 21 24 11057 0 0 1 0 B718603 1 1 1 21 11058 0 0 1 0 B718636 22 30 4 11058 0	nent	Prescreen No.	YF	VE	H	AVS No.	VF	JE	VE	FF	SF	W
B718603 1 1 21 11058 0 0 1 0 B718636 22 30 4 11059 0 0 0 0 B718648 25 16 27 11060 1 0 0 0 0 B718649 18 7 3 11061 0	P	B718599	0	21	24	11057	0	0	1	0	0	IN
B718636 22 30 4 11059 0 <	14P	B718603	1	1	21	11058	0	0	1	0	0	NT
B718648 25 16 27 11060 1 0 2 1 B718649 18 7 3 11061 0 <td>14P</td> <td>B718636</td> <td>22</td> <td>30</td> <td>4</td> <td>11059</td> <td>0</td> <td>0</td> <td>0</td> <td>0</td> <td>1</td> <td>NT</td>	14P	B718636	22	30	4	11059	0	0	0	0	1	NT
B718649 18 7 3 11061 0 0 0 0 B718656 22 7 58 11062 z1 0 <td>IP</td> <td>B718648</td> <td>25</td> <td>91</td> <td>72</td> <td>11060</td> <td>1</td> <td>0</td> <td>2</td> <td>1</td> <td>7</td> <td>NT</td>	IP	B718648	25	91	72	11060	1	0	2	1	7	NT
B718656 22 7 58 11062 z1 0	IP	B718649	18	7	3	11061	0	0	0	0	0	NT
B718789 30 19 0 11063 0 <	IP	B718656	22	7	28	11062	z1	0	0	0	7	N
B718799 17 32 5 11064 0 0 2 0 B718807 7 19 36 11065 0 0 1 0 B718813 0 46 6 11066 0 0 1 0 B718819 0 38 23 11067 0 0 1 0 B718821 0 8 24 11068 0 0 1 0 B718832 6 32 31 11070 0 0 1 0 B718840 1 20 13 11071 0 0 0 0 0 B718842 10 14 21 11072 0 0 0 0 0 B718843 39 11 0 11073 1 0 0 0 0 0 B718860 38 13 0 11075 1 0 <td>IP</td> <td>B718789</td> <td>30</td> <td>61</td> <td>0</td> <td>11063</td> <td>0</td> <td>0</td> <td>0</td> <td>0</td> <td>0</td> <td>NT</td>	IP	B718789	30	61	0	11063	0	0	0	0	0	NT
B718807 7 19 36 11065 0 0 1 0 B718813 0 46 6 11066 0 1 0 1 0 1 0 0 1 0 0 1 0 0 1 0 0 1 0 0 1 0 <td>IP</td> <td>B718799</td> <td>11</td> <td>32</td> <td>2</td> <td>11064</td> <td>0</td> <td>0</td> <td>2</td> <td>0</td> <td>0</td> <td>N</td>	IP	B718799	11	32	2	11064	0	0	2	0	0	N
B718813 0 46 6 11066 0 0 1 0 B718819 0 38 23 11067 0 0 1 0 B718821 0 8 24 11068 0 0 1 0 B718842 1 20 13 11070 0 0 0 0 0 B718843 39 11 0 11073 0 0 0 0 0 0 B718849 7 18 25 11074 0 0 0 0 0 B718860 38 13 0 11075 1 0 1 0 1 B718862 4 7 16 11076 0 0 0 0 0 0 B718876 33 1077 0 0 0 0 0 0 0 0 B718876 4	IP	B718807	7	61	36	11065	0	0	1	0	1	N
B718819 0 38 23 11067 0 0 1 0 B718831 6 32 31 11069 0 0 1 0 B718838 6 32 31 11069 0 0 1 0 B718840 1 20 13 11071 0 0 0 0 0 B718842 10 14 21 11072 0 0 1 0 0 B718843 39 11 0 11073 1 0 1 0 0 0 B718849 7 18 25 11074 0 0 0 0 2 B718860 38 13 0 11075 1 0 0 0 0 0 B718876 4 7 16 1077 0 0 0 0 0 0 0	IP	B718813	0	46	9	11066	0	0	1	0	0	NT
B718821 0 8 24 11068 0 0 1 0 B718838 6 32 31 11069 0 0 1 0 B718840 1 20 13 11070 0 0 0 0 0 B718841 7 44 33 11071 0 0 0 0 0 0 B718842 10 14 21 11072 0 0 1 0	IP	B718819	0	38	23	11067	0	0	1	0	0	NT
B718838 6 32 31 11069 0 0 1 0 B718840 1 20 13 11070 0 0 0 0 B718841 7 44 33 11071 0 0 0 0 0 B718842 10 14 21 11072 0 0 1 0 <t< td=""><td>IP.</td><td>B718821</td><td>0</td><td>80</td><td>24</td><td>11068</td><td>0</td><td>0</td><td>1</td><td>0</td><td>0</td><td>Į.</td></t<>	IP.	B718821	0	80	24	11068	0	0	1	0	0	Į.
B718840 1 20 13 11070 0 0 0 0 B718841 7 44 33 11071 0 0 0 0 0 B718843 39 11 0 11073 1 0 1 0 1 0 1 0	IP.	B718838	9	32	31	11069	0	0	1	0	0	¥
B718841 7 44 33 11071 0 0 0 0 B718842 10 14 21 11072 0 0 1 0 B718843 39 11 0 11073 1 0 1 0 1 B718849 7 18 25 11074 0 0 0 0 2 B718860 38 13 0 11075 1 0 1 1 1 B718862 4 7 16 11076 1 0 0 0 0	P	B718840	1	20	13	11070	0	0	0	0	1	¥
B718842 10 14 21 11072 0 0 1 0 B718843 39 11 0 11073 1 0 1 0 </td <td>IP.</td> <td>B718841</td> <td>7</td> <td>44</td> <td>33</td> <td>11071</td> <td>0</td> <td>0</td> <td>0</td> <td>0</td> <td>0</td> <td>FN</td>	IP.	B718841	7	44	33	11071	0	0	0	0	0	FN
B718843 39 11 0 11073 1 0 1 0 B718849 7 18 25 11074 0 0 0 2 B718860 38 13 0 11075 1 0 1 1 B718862 4 7 16 11076 1 0 0 0 0 0	IP.	B718842	10	14	21	11072	0	0	1	0	0	TN.
B718849 7 18 25 11074 0 0 0 2 B718860 38 13 0 11075 1 0 1 <td>F</td> <td>B718843</td> <td>39</td> <td>11</td> <td>0</td> <td>11073</td> <td>1</td> <td>0</td> <td>1</td> <td>0</td> <td>0</td> <td>N</td>	F	B718843	39	11	0	11073	1	0	1	0	0	N
B718860 38 13 0 11075 1 0 1 1 B718862 4 7 16 11076 1 0 0 2 B718876 53 6 0 11077 0 0 0 0	IP II	B718849	7	18	25	11074	0	0	0	2	0	IN
B718862 4 7 16 11076 1 0 0 2 B718876 53 6 0 11077 0	IP.	B718860	38	13	0	11075	1	0	1	1	0	IN
B718876 53 6 0 11077 0 0 0 0 0	Ь	B718862	4	7	16	11076	1	0	0	2	1	Ā
	14P	B718876	53	9	0	11077	0	0	0	0	0	Į.

Results of Prescreen Testing	
Results of Prescreen	Testinga
Results of	Prescreen
Result	s of
	Result

W		IN	N.	M	N	NT	IN	NT	TN	IN	TN	TN	N.	ŢN	NT	IN	IN	IN	Į.	Į.	0	0
JE VE PT SF		0	0	0	0	0	0	0	0	1	0	0	0	0	0	0	0	0	1	0	0	zl
R		0	0	0	0	0	0	0	0	2	1	1	0	1	1	0	0	1	1	1	0	0
VE		2	1	1	1	1	1	1	1	3	1	1	1	1	0	0	2	1	2	1	0	1
JE		0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
YF		0	0	0	0	0	0	0	0	0	0	0	0	1	0	0	0	0	1	1	0	1
AVS	No.	11078	11079	11080	11081	11082	11083	11084	11085	11086	11087	11088	11089	11090	11091	11092	11093	11094	11095	11096	9131	9132
F		3	0	13	8	16	14	10	80	7	1	48	21	13	20	13	19	38	53	14	8	15
VE		41	22	39	31	28	16	21	25	62	17	26	45	16	13	7	23	47	25	13	1	10
YF		1	1	5	9	27	10	1	2	4	4	9	0	5	4	2	9	3	25	22	26	16
Prescreen	No.	B715882	B718889	B718917	B718924	B718934	B718935	B718942	B718954	B718969	B718978	B718979	B718982	B718983	B718990	B718996	B719004	B719006	B719007	B719008	B719215	B719216
Shipment	No.	14P	12P	12P																		

Results of Prescreen Testing

		este e Baanta	75			PIPLES!	5.11W-15-	E.,						21. ·							
M	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	1	0	0	0	0
SF	0	1	0	0	0	1	0	0	0	1	0	0	0	0	0	0	0	1	0	1	0
FT	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	2	1	. 1	1	2
VE	0	2	1	0	1	1	0	0	0	0	0	0	1	0	0	1	1	1	0	1	1
JE	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
YF	0	1	1	0	0	0	0	0	0	0	0	0	0	0	0	0	1	1	0	1	0
AVS No.	9133	9134	9135	9136	9137	9138	9139	9140	9141	9142	9143	9144	9145	9146	9147	9148	9149	9150	1516	9152	9153
m	2	11	9	13	10	32	13	5	7	3	6	11	36	28	8	15	37	8	16	16	18
VE	1	64	7	2	27	6	1	1	3	9	23	9	10	4	15	2	15	15	9	3	9
YF	16	33	33	22	16	19	28	27	13	13	111	6	7	10	14	14	20	13	91	20	16
Prescreen No.	B719217	B719218	B719219	B719220	B719221	B719222	B719224	B719225	B719226	B719228	B719229	B719230	B719232	B719238	B719240	B719241	B719242	B719244	B719245	B719246	B719247
Shipment No.	12P	12P	12P	12P	12P	12P	12P	12P	12P	12P	12P	12P	12P	12P	12P	12P	12P	12P	12P	12P	12P

Results of Prescreen Testinga

	3	0	0	0	0	0	0	0	0	0	0	0	0	FN	M	FN	F	FN	F	Ę	F	F
	SF	0	0	0	0	0	0	0	0	0	0	0	0	0	1	1	0	0	0	0	0	0
	Z	0	0	0	0	0	0	1	0	0	0	1	0	0	0	0	0	0	1	0	1	0
	VE	0	0	0	0	0	0	1	1	2	2	0	1	2	1	0	3	2	2	2	0	0
	JE	0	0	0	0	0.	0	0	0	0	0	0	0	0	0	0	0	2	0	0	0	0
F	YF	0	0	0	1	0	0	1	1	0	1	2	0	0	0	0	0	0	0	0	0	0
	AVS No.	9154	9155	9156	9157	9158	9159	0916	1916	9162	9163	9164	9165	11097	11098	11099	11100	11101	11102	11103	11104	11105
	E	14	17	4	2	5	19	4	11	19	5	5	10	30	0	0	24	11	0	12	21	0
	VE	1	9	3	2	5	9	16	36	30	20	1	10	72	1	19	72	52	4	22	2	4
1	E	10	5	13	16	23	15	6	7	10	8	18	0	5	21	0	2	3	23	0	33	20
	Prescreen No.	B719255	B719256	B719257	B719258	B719259	B719260	B719265	B719266	B719267	B719272	B719274	B719278	B720392	B720394	B720396	B720398	B720399	B720400	B720405	B720409	B720410
	Shipment No.	12P	14P																			

Results of Prescreen Testing

Results of Primary Screen Testingb

Mo. AVS XVB FT AVS XVB FT SSP VVB NO	The same		Kesults	S of Fresci	of Frescreen Testing	The Line of the leading of the lead	20010000	Kesults	Results of Primary Screen Testing	v Screen I	esting	
B720412 42 5 13 11106 0 0 1 0 B720415 32 3 1 11107 0 0 1 2 0 B720418 4 8 22 11108 0 0 2 0 1 B720418 21 12 0 11109 0 0 2 0 1 B720425 41 32 1 11109 0	Shipment No.	1,000,000		VE	PT	AVS No.	YF	æ	VE	ΙĀ	SF	W
B720415 32 3 1 11107 0 0 1 2 0 B720418 4 8 22 11108 0 0 0 2 0 B720419 21 12 0 11109 0 0 2 0 1 B720424 9 5 27 11110 0 0 2 0 1 B720425 41 32 1 11111 0 0 2 0	14P	B720412	42	5	13	11106	0	0	0	1	0	Ţ
B720418 4 8 22 11108 0 0 0 2 0 B720419 21 112 0 11109 0 0 2 0 1 B720424 9 5 27 11110 0 0 2 0 1 B720425 41 32 1 11111 0 0 2 0 1 B720427 - - - - 11111 NT NT NT NT B720430 13 12 51 11115 0 0 1 0 0 B720430 - - - - 11114 NT NT NT NT B720430 - - - 11115 NT NT NT NT B720430 - - - - 11115 NT NT NT NT B720431 - -	14P	B720415	32	3	1	11107	0	0	1	7	0	¥
B720419 21 12 0 11109 0 2 0 1 B720424 9 5 27 11110 0 0 2 0 1 B720425 41 32 1 11111 0 0 2 0	14P	B720418	4	8	22	11108	0	0	0	2	0	K
B720424 9 5 27 1111 0 0 2 0 B720425 41 32 1 1111 0 0 2 0 0 B720425 -1 -1 1111 NT NT NT NT NT B720427 -2 -2 1111 NT NT NT NT NT B720430 13 12 51 11115 0 0 1 0	14P	B720419	21	12	0	11109	0	0	2	0	1	N.
B720425 41 32 1 11111 0 0 2 0 0 B720425 - - - - 11112 NT NT NT NT B720427 - - - 11113 0 0 1 0 0 B720427 - - - 11114 NT NT NT NT B720430 13 12 51 11115 0 1 2 0 B720430 - - - 11116 NT NT NT NT B720431 14 11 8 11117 0 1 2 1 0 B720433 28 6 3 11118 1 0 1 2 1 0 B720435 5 20 10 11120 0 0 0 0 0 0 0 0 0 0	14P	B720424	6	5	27	11110	0	0	0	2	0	M
B720425 - - - - - - - NT NT <td>14P</td> <td>B720425</td> <td>41</td> <td>32</td> <td>1</td> <td>11111</td> <td>0</td> <td>0</td> <td>2</td> <td>0</td> <td>0</td> <td>F</td>	14P	B720425	41	32	1	11111	0	0	2	0	0	F
B720427 25 28 1 11113 0 0 1 0 0 B720427 - - - - - 11114 NT NT NT NT B720430 13 12 51 11115 0 0 1 2 0 B720430 - - - - 11116 NT NT NT NT B720431 14 11 8 11117 0 0 2 1 0 B720433 28 6 3 11118 1 0 1 2 1 0 B720434 8 19 0 11120 0 <t< td=""><td>14P</td><td>B720425</td><td>1</td><td></td><td>1</td><td>11112</td><td>NT</td><td>NT</td><td>NT</td><td>IN</td><td>IN</td><td>NT</td></t<>	14P	B720425	1		1	11112	NT	NT	NT	IN	IN	NT
B720427 - </td <td>14P</td> <td>B720427</td> <td>25</td> <td>28</td> <td>1</td> <td>11113</td> <td>0</td> <td>0</td> <td>1</td> <td>0</td> <td>0</td> <td>N.</td>	14P	B720427	25	28	1	11113	0	0	1	0	0	N.
B720430 13 12 51 11115 0 0 1 2 0 B720430 11116 NT NT NT NT B720431 14 11 8 11117 0 0 2 1 0 B720433 28 6 3 11118 1 0 1 2 1 0 B720435 5 20 10 11120 0	14P	B720427	-	-	1	11114	NT	NT	NT	TN	NT	IN
B720430 - - - - - - - - - - NT NT NT NT NT NT B720431 14 11 8 11117 0 0 2 1 0 1 0 1 0 1 0 <td>14P</td> <td>B720430</td> <td>13</td> <td>12</td> <td>51</td> <td>11115</td> <td>0</td> <td>0</td> <td>1</td> <td>2</td> <td>0</td> <td>TN.</td>	14P	B720430	13	12	51	11115	0	0	1	2	0	TN.
B720431 14 11 8 11117 0 0 2 1 0 B720433 28 6 3 11118 1 0 1 2 1 B720434 8 19 0 11119 0 0 0 0 0 0 B720435 5 20 10 11120 0	14P	B720430	1	-	ŀ	11116	NT	NT	NT	NT	IN	N.
B720433 28 6 3 11118 1 0 1 2 1 B720434 8 19 0 11119 0 0 2 0 0 B720435 5 20 10 11120 0 0 0 0 0 0 B720436 28 4 0 11121 0 0 2 3 1 8 B720443 25 1 5 11122 1 2 0 2 2 2 B720445 42 4 82 11123 2 1 3 2 2 B720490 19 3 3 8507 0 0 0 0 0 0 0 B720493 26 4 1 8509 0 0 0 0 0 0 0	14P	B720431	14	11	••	11117	0	0	2	1	0	N.
B720434 8 19 0 11119 0 2 0 0 B720435 5 20 10 11120 0 0 0 0 0 B720436 28 4 0 11121 0 0 2 3 1 B720443 25 1 5 11122 1 2 0 2 3 1 B720445 42 4 82 11123 2 1 3 2 B720490 19 3 3 8507 0 0 0 0 0 0 B720493 26 4 1 8509 0 0 0 0 0 0 0	14P	B720433	28	9	3	11118	1	0	1	2	1	K
B720435 5 20 10 11120 <	14P	B720434	8	61	0	11119	0	0	2	0	0	M
B720436 28 4 0 11121 0 0 2 3 1 B720443 25 1 5 11122 1 2 0 2 2 B720445 42 4 82 11123 2 1 1 3 2 B720490 19 3 3 8507 0 0 0 0 0 B720492 20 4 1 8509 0 0 0 0 0 0	14P	B720435	2	20	10	11120	0	0	0	0	0	NT
B720443 25 1 5 11122 1 2 0 2 2 B720445 42 4 82 11123 2 1 1 3 2 B720490 19 3 3 8507 0 0 0 0 0 B720492 20 4 1 8509 0 0 0 0 0 0	14P	B720436	28	4	0	11121	0	0	2	3	1	NT
B720445 42 4 82 11123 2 1 1 3 2 B720490 19 3 3 8507 0	14P	B720443	25	1	2	11122	1	2	0	2	2	TN.
B720490 19 3 3 8507 0 <th< td=""><td>14P</td><td>B720445</td><td>42</td><td>4</td><td>82</td><td>11123</td><td>2</td><td>1</td><td>1</td><td>3</td><td>2</td><td>M</td></th<>	14P	B720445	42	4	82	11123	2	1	1	3	2	M
B720492 20 0 2 8508 0 0 0 0 0 0 B720493 26 4 1 8509 0 0 0 0 0 0	8P	B720490	19	3	3	8507	0	0	0	0	0	0
B720493 26 4 1 8509 0 0 0 0 0 0	8P	B720492	20	0	2	8208	0	0	0	0	0	1
	8P	B720493	26	4	1	8509	0	0	0	0	0	0

Results of Prescreen Testing*

ent Prescreen YF VE PT AVS YF JE B720494 18 20 3 8273 0 0 B720498 17 8 2 8510 0 0 B720420 17 12 5 9166 1 0 0 B720420 16 1 4 9167 1 0 0 B720422 14 18 50 9168 1 0 0 B720423 0 13 27 9170 0 0 0 B720432 0 13 27 9170 0 0 0 B720432 0 13 27 9171 0 0 0 B720434 18 - 27 7324 0 0 0 B720483 18 - 29 7326 0 0 0 B7204895 18 -					9							
B720494 18 20 3 8273 0 0 B720498 17 8 2 8510 0 0 B720498 17 12 5 9166 1 0 B720821 16 1 4 9167 1 0 B720822 14 18 50 9168 1 0 B720832 14 18 50 9169 1 0 B720832 0 13 27 9170 0 0 B720832 0 13 27 9171 0 0 B720834 7 31 25 9171 0 0 B720835 18 - 27 7324 0 0 0 B720896 2 - 29 7326 0 0 0 B720900 28 - 41 7328 1,1 0 0 B720910 </th <th>Shipment No.</th> <th>Prescreen No.</th> <th>VF</th> <th>VE</th> <th>T.</th> <th>AVS No.</th> <th>YF</th> <th>JE</th> <th>VE</th> <th>12</th> <th>SF</th> <th>A</th>	Shipment No.	Prescreen No.	VF	VE	T.	AVS No.	YF	JE	VE	1 2	SF	A
B720498 17 8 2 8510 0 0 B720820 7 12 5 9166 1 0 B720821 16 1 4 9167 1 0 B720825 14 18 50 9168 1 0 B720825 14 18 50 9169 1 0 B720836 23 9 1 9169 1 0 B720837 0 13 27 9170 0 0 B720834 7 31 25 9171 0 0 B720894 18 - 27 7324 0 0 0 B720895 18 - 27 7326 0 0 0 B720896 22 - 29 7326 0 0 0 B720903 5 - 29 7329 0 0 0	8P	B720494	18	20	3	8273	0	0	0	0	0	0
B720820 7 12 5 9166 1 0 B720821 16 1 4 9167 1 0 B720825 14 18 50 9168 1 0 B720826 23 9 1 9169 1 0 B720832 0 13 27 9170 0 0 B720834 7 31 25 9171 0 0 B720834 7 31 25 9171 0 0 B720895 6 - 27 7324 0 0 B720896 22 - 29 7325 0 0 B720896 22 - 35 7326 0 0 B720900 28 - 45 7329 0 0 B720901 23 11 22 8511 2,2 1,0 B720911 14 14	8P	B720498	17	8	2	8510	0	0	1	0	0	0
B720821 16 1 4 9167 1 0 B720828 14 18 50 9168 1 0 B720828 23 9 1 9169 1 0 B720832 0 13 27 9170 0 0 B720834 7 31 25 9171 0 0 B720895 6 - 27 7324 0 0 B720895 18 - 29 7326 0 0 B720896 22 - 35 7326 0 0 B720896 22 - 35 7326 0 0 B720896 22 - 29 7326 0 0 B720900 28 - 45 7330 0 0 B720910 23 11 22 8511 2,2 1,0 B720912 6 - 4	12P	B720820	7	12	5	9916	1	0	2	0	0	1
B720825 14 18 50 9168 1 0 B720828 23 9 1 9169 1 0 B720832 0 13 27 9170 0 0 B720834 7 31 25 9171 0 0 B720895 18 - 27 7324 0 0 0 B720896 18 - 29 7325 0 0 0 B720896 22 - 29 7326 0 0 0 B720896 22 - 36 7326 0 0 0 B720900 28 - 41 7328 1,1 0,0 0 B720903 5 - 29 7329 0 0 0 B720910 23 11 22 8511 2,2 1,0 0 B720912 6 - 43 7332 </td <td>12P</td> <td>B720821</td> <td>16</td> <td>1</td> <td>4</td> <td>2916</td> <td>1</td> <td>0</td> <td>1</td> <td>2</td> <td>0</td> <td>0</td>	12P	B720821	16	1	4	2916	1	0	1	2	0	0
B720828 23 9 1 9169 1 0 B720832 0 13 27 9170 0 0 B720834 7 31 25 9171 0 0 B720892 6 - 27 7324 0 0 B720894 18 - 29 7325 0 0 B720895 18 - 29 7326 0 0 B720896 22 - 30 7327 0 0 B720900 28 - 41 7328 1,1 0,0 B720903 5 - 29 7329 0 0 0 B720904 6 - 45 7330 0 0 0 B720911 14 14 55 8511 2,2 1,0 0 B720912 6 - 43 7331 0,0 0,0 B7	12P	B720825	14	18	50	8916	1	0	1	1	0	0
B720832 0 13 27 9170 0 0 B720834 7 31 25 9171 0 0 B720892 6 - 27 7324 0 0 B720894 18 - 29 7325 0 0 B720895 18 - 35 7326 0 0 B720806 22 - 30 7327 0 0 B720900 28 - 41 7328 1,1 0,0 B720905 6 - 45 7329 0 0 B720910 14 14 55 8511 2,2 1,0 B720911 14 14 55 8272 1,0 0,0 B720914 33 - 43 7331 0,0 0,0 B720915 4 - 37 7332 2,1 0,0 B720915 4 - <td>12P</td> <td>B720828</td> <td>23</td> <td>6</td> <td>1</td> <td>6916</td> <td>1</td> <td>0</td> <td>1</td> <td>0</td> <td>0</td> <td>0</td>	12P	B720828	23	6	1	6916	1	0	1	0	0	0
B720834 7 31 25 9171 0 0 B720892 6 - 27 7324 0 0 B720894 18 - 29 7325 0 0 B720895 18 - 35 7326 0 0 B720896 22 - 30 7327 0 0 B720900 28 - 41 7328 1,1 0,0 B720903 5 - 29 7330 0 0 B720910 23 11 22 8511 2,2 1,0 B720911 14 14 55 8511 2,2 1,0 B720914 33 - 43 7331 0,0 0,0 B720915 4 - 31 7333 0,0 0,0 B720915 4 - 31 7333 0,0 0,0	12P	B720832	0	13	27	9170	0	0	1	0	0	0
B720892 6 - 27 7324 0 0 B720894 18 - 29 7325 0 0 B720895 18 - 35 7326 0 0 B720896 22 - 30 7327 0 0 B720900 28 - 41 7328 1,1 0,0 B720903 5 - 29 7329 0 0 B720910 23 11 22 8511 2,2 1,0 B720911 14 55 8512 1,0 0,0 B720912 6 - 43 7331 0,0 0,0 B720914 33 - 37 7332 2,1 0,0 B720915 4 - 31 7333 0,0 0,0 B720915 4 - 31 7333 0,0 0,0	12P	B720834	7	31	25	1716	0	0	2	0	0	0
B720894 18 29 7325 0 0 B720895 18 35 7326 0 0 B720806 22 30 7327 0 0 B720900 28 41 7328 1,1 0,0 B720903 5 29 7329 0 0 0 B720905 6 45 7330 0 0 0 B720910 23 11 22 8511 2,2 1,0 0 B720912 6 43 7331 0,0 0,0 0 B720914 33 43 7332 2,1 0,0 0 B720915 4 31 7333 0,0 0,0 0	4P	B720892	9	-	27	7324	0	0	0	0	0	0
B720895 18 - 35 7326 0 0 B720896 22 - 30 7327 0 0 B720900 28 - 41 7328 1,1 0,0 B720903 5 - 29 7329 0 0 B720905 6 - 45 7330 0 0 B720910 23 11 22 8511 2,2 1,0 B720911 14 14 55 8272 1,0 0,0 B720912 6 - 43 7331 0,0 0,0 B720915 4 - 37 7332 2,1 0,0 B720915 4 - 31 7333 0,0 0,0	4P	B720894	18	1	29	7325	0	0	0	0	0	0
B720896 22 — 30 7327 0 0 B720900 28 — 41 7328 1,1 0,0 B720903 5 — 29 7329 0 0 B720905 6 — 45 7330 0 0 B720910 23 11 22 8511 2,2 1,0 B720911 14 14 55 8212 1,0 0,0 B720912 6 — 43 7331 0,0 0,0 B720915 4 — 37 7332 2,1 0,0 B720915 4 — 31 7333 0,0 0,0	4P	B720895	18	1	35	7326	0	0	0	0	0	0
B720900 28 - 41 7328 1,1 0,0 B720903 5 - 29 7329 0 0 B720905 6 - 45 7330 0 0 B720910 23 11 22 8511 2,2 1,0 B720911 14 14 55 8272 1,0 0,0 B720912 6 - 43 7331 0,0 0,0 B720914 33 - 37 7332 2,1 0,0 B720915 4 - 31 7333 0,0 0,0	4P	B720896	22	1	30	7327	0	0	0	0	0	0
B720903 5 - 29 7329 0 0 B720905 6 - 45 7330 0 0 0 B720910 23 11 22 8511 2,2 1,0 0 B720911 14 14 55 8272 1,0 0,0 0 B720912 6 - 43 7331 0,0 0,0 0 B720914 33 - 37 7332 2,1 0,0 0 B720915 4 - 31 7333 0,0 0,0 0	4P	B720900	28	-	41	7328	1,1	0,0	0,0	0,0	0,0	0
B720905 6 - 45 7330 0 0 B720910 23 11 22 8511 2,2 1,0 B720911 14 14 55 8272 1,0 0,0 B720912 6 - 43 7331 0,0 0,0 B720914 33 - 37 7332 2,1 0,0 B720915 4 - 31 7333 0,0 0,0	4P	B720903	5	1	29	7329	0	0	0	0	0	0
B720910 23 11 22 8511 2,2 1,0 B720911 14 14 55 8272 1,0 0,0 B720912 6 - 43 7331 0,0 0,0 B720914 33 - 37 7332 2,1 0,0 B720915 4 - 31 7333 0,0 0,0	4P	B720905	9	1	45	7330	0	0	0	0	0	0
B720911 14 14 55 8272 1,0 0,0 B720912 6 - 43 7331 0,0 0,0 B720914 33 - 37 7332 2,1 0,0 B720915 4 - 31 7333 0,0 0,0	8P	B720910	23	11	22	8511	2,2	1,0	2,1	2,2	2,2	0
B720912 6 - 43 7331 0,0 0,0 B720914 33 - 37 7332 2,1 0,0 B720915 4 - 31 7333 0,0 0,0	8P	B720911	14	14	55	8272	1,0	0,0	1,0	2,0	1,0	0
B720914 33 - 37 7332 2,1 0,0 B720915 4 - 31 7333 0,0 0,0	4P	B720912	9	-	43	7331	0,0	0,0	1,0	0,0	1,1	0
B720915 4 - 31 7333 0,0 0,0	4P	B720914	33	-	37	7332	2,1	0,0	1,1	0,1	2,2	0
2000000	4P	B720915	4	-	31	7333	0,0	0,0	0,0	0,0	2,2	0
B720910 9 - 55 7099 0 0	4P	B720916	6	1	55	7099	0	0	0	0	0	0

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Testing
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Prescreen
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Results

		Results		of Prescreen Testing ^a	M.		Results	Results of Primary Screen Testing ^b	y Screen T	esting	
Shipment No.	Prescreen No.	YF	VE	L	AVS No.	YF	æ	VE	FT	SF	A
8P	B720931	0	25	46	8274	0	0	0	1	1	0
8P	B720933	0	100	47	8513	0,1	0,0	3,3	0,1	1,1	-
8P	B720935	18	10	2	8325	0	0	0	0	0	0
4P	B720937	111	1	55	7100	0	0	0	0	0	•
4P	B720940	2	1	09	7101	0	0	0	0	0	•
4P	B720941	0	1	51	7102	0	0	0	0	0	0
4P	B720946	9	1	32	7336	0	0	1	0	1	0
4P	B720951	11	1	62	7103	0	0	0	0	0	2
4P	B720951	-	-	•	7337	NT	NT	NT	IN	NT	IN.
4P	B720952	11	1	99	7104	0	0	0	0	0	0
4P	B720952	1	1	1	7338	IN	NT	NT	NT	NT	N
4P	B720958	1	1	09	7105	0	0	0	0	0	0
4P	B720963	0	1	59	7106	0	0	0	0	1	1
8P	B720968	0	26	40	8514	1	0	1	1	0	0
4P	B720978	10	-	24	7339	1,0	1,0	1,0	0,0	0,1	0
4P	B720979	10	_	63	7107	0	0	0	0	1	0
4P	B720979	1	-	1	7340	TN	NT	NT	M	TN	K
4P	B720985	7	-	0	7341	1,1	0,0	0,1	0,1	0,0	0
4P	B720986	7	-	0	7342	0	0	0	0	0	0
4P	B720987	12	1	17	7343	0	0	1	0	1	•
4P	B720988	4	1	4	7344	1,0	0,0	1,0	0,0	2,0	0

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Testing
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rescr
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Results

Shipment No.	Prescreen No.	YF	VE	P	AVS No.	YF	JE	VE	P	SF	M
	B720989	28	1	1	7345	0	0	0	0	0	0
	B720990	7	1	23	7346	0	0	0	0	0	0
	B720994	4	1	0	7347	0,0	0,0	1,0	0,0	1,0	0
	B720995	7	1	0	7348	0	0	0	0	0	0
	B720998	8	1	1	7349	0,0	0,0	0,0	2,1	0,0	0
	B721000	12	1	24	7350	1,1	0,1	0,0	2,0	1,0	0
	B721001	3	1	3	7351	1,1	0,0	0,0	0,0	0,0	0
	B721002	2	1	27	7352	0	0	0	0	0	0
	B721005	10	1	12	7353	0	0	0	0	0	0
	B721006	32	1	2	7354	2,2	0,1	0,0	0,0	0,0	0
	B721007	36	1	80	7355	0,0	0,0	0,0	1,0	0,0	0
	B721011	37	-	09	7298	0,0	0,0	1,0	0,0	0,0	1
	B721012	32	-	36	7356	1,0	0,0	0,0	1,0	2,0	0
	B721013	0	1	10	7357	0	0	0	0	0	0
	B721015	7	1	0	7358	1,0	0,0	0,0	0,0	1,1	0
- 7	B721016	0	1	34	7359	0	0	0	0	0	0
	B721017	0	1	39	7360	0	0	0	0	0	0
	B721018	3	-	32	7361	0,0	0,0	1,0	0,0	0,0	0
	B721019	0	1	28	7362	0,0	0,0	0,0	1,0	0,0	0
	B721020	4	1	10	7363	1,0	1,0	1,0	0,0	1,0	0
	B721021	0	ı	61	7299	0	0	0	0	0	1

Results of Prescreen Testing

Results of Primary Screen Testing	PT SF VV	1 0 0	0,0 2,0 0	0 0 0	1 0 0	2,0 0,0 0	0,0 2,0 0	0 0 0	0 0 1	0 0 0	0,0 1,0 0	2,2 2,1 0	1,1 0,0 1,1	2,1 1,0 2	0 0 0	0,0 1,0 0	0,0 2,0 1	1,1 0,0 0	0,0 1,0 0	0,1 0,0 1	1,0 2,1 0	
of Primary S	VE	0	0,0	0	0	0,0	0,0	0	0	0	0,0	0,0	0,0	0,0	0	0,0	1,0	0,0	0,0	0,1	0,0	
Kesults	æ	0	0,0	0	0	0,0	0,0	0	0	0	1,1	0,0	1,0	2,0	0	1,0	0,0	0,0	1,0	1,0	0,0	
	YF	0	0,1	0 9	0 4	0,0	0,0	0 (5 1	0	0,1	3 2,1	0,0	5 2,0	0 9	2,0	2,1	0,0	0,1	0,0	0,1	
sting"	r Avs	7364	7365	7366	7367	3 7368	7369	7370	8515	7371	7372	7373	7374	375	7376	T377	5 7378	7379	7380	7381	7382	
is of Prescreen Testing	VE PT	- 32	- 37	- 29	- 25	- 43	_ 2	- 10	28 22	- 5	0	- 17		- 15	4	0 -	- 15	7	_ 20	0	- 4	
Results of	YF	0	3	0	12	0	10	11	0	7	8	13	10	47	12	•	35	12	10	14	19	
	Prescreen No.	B721022	B721026	B721029	B721030	B721031	B721033	B721034	B721035	B721040	B721042	B721045	B721050	B721051	B721052	B721053	B721054	B721055	B721056	B721058	B721059	
	Shipment No.	4P	4P	4P	4P	4P	4P	4P	8P	4P	4P	4P	4P	4P	4P	4P	4P	4P	4P	4P	4P	

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		Result	s of Prescr	is of Prescreen Testing			Results	Results of Primary Screen Testing ^b	y Screen T	esting	
Shipment No.	Prescreen No.	YF	VE	M	AVS No.	YF	JE	VE	M	SF	W
4P	B721061	1	-	81	7301	0,0	0,0	0'0	1,2	0,1	0
46	B721062	18	1	86	7302	0,0	0,0	0,0	0,1	1,1	2,1
4P	B721063	15	1	36	7383	1,2	0,0	0,0	2,0	1,1	0
4P	B721064	8	1	80	7303	0,0	0,0	0,1	1,2	0,0	0
4P	B721173	62	-	20	7108	1,1	1,0	0,0	0,0	0,0	0
4P	B721177	10	-	6	7385	1,0	0,0	0,1	2,0	1,0	0
4P	B721178	3	-	10	7386	0,1	0,0	0,2	2,1	0,0	0
8P	B721181	20	10	4	8326	2,1	0,0	0,0	1,1	2,2	0
8P	B721192	21	S	1	8327	2,1	0,0	0,1	1,0	0,0	0
4P	B721193	0	-	23	7387	0,0	0,0	0,0	1,0	0,0	0
4P	B721196	-	-	31	7388	0	0	0	0	0	0
4P	B721211	27	1	0	8334	0	0	0	0	0	0
8P	B721213	15	3	2	8504	0	0	0	0	0	0
8P	B721220	24	8	3	8505	NT	NT	1	0	0	0
8P	B721234	18	8	0	8506	INT	NT	1	1	1	1
8P	B721236	15	22	17	8328	0	0	1	2	1	0
4P	B721256	14	-	36	7389	0	0	0	0	0	0
4P	B721257	12	-	10	7390	2,0	0,0	0,0	0,1	1,0	0
8P	B721258	18	3	11	8329	0	0	0	0	0	0
4P	B721260	10	1	21	7391	0,0	0,0	0,0	2,1	1,1	2,1
4P	B721261	8	1	19	7392	0	0	0	0	-	0

Results of Prescreen Testing

VE	1000
9	9 -
. 12	- 12
2	4 2
09	09 9
2 40	12 40
. 39	- 39
. 38	- 38
24	3 24
26	4 56
. 61	- 61
49	- 64
. 51	- 51
. 52	- 52
48	2 48
37	2 37
41	2 41
42	2 42
3 56	23 56
35	0 35
7 29	27 29
3 46	43 46

	W	0	0	0	2	0	0	0	1	0	0	0	0	1	0	2	-	0	0	0	0	0
Sting	SF	0	0	1	0	0,0	0	0	0,0	0	0	1,0	1	1	0	0	0,0	0	0	0	1,1	0
Screen Te	M	1	0	0	0	0,2	0	0	0,0	1	1	1,1	0	1	0	0	1,0	0	0	0	2,2	0
Results of Primary Screen Testing	VE	0	0	0	1	1,1	0	0	1,0	0	0	0,0	0	1	0	0	0,0	1	0	0	0,1	0
Kesults o	E	0	1	0	1	0,0	0	0	0,0	0	0	0,0	0	0	0	0	0,0	0	0	0	0,0	0
	ž.	1	1	1	1	0,0	0	0	0,0	0	0	1,1	0	0	0	0	1,0	1	0	0	1,1	0
	AVS No.	8319	8335	8336	8337	7308	8320	7309	7310	8321	8322	7395	8323	8500	8501	7396	7311	8502	7397	7398	7399	7400
esults of Prescreen Testing	m .	30	€	0	2	59	35	55	52	27	38	44	72	52	33	22	50	37	44	24	32	45
s of Prescr	VE	2	1	1	-	1	9	1	1	1	1	1	2	14	0	1	1	6	1	•	1	1
Kesult	YF	3	40	32	35	28	1	14	11	9	0	12	18	0	0	5	10	0	7	5	34	6
	Prescreen No.	B721582	B721584	B721586	B721588	B721592	B721594	B721595	B721596	B721598	B721601	B721602	B721604	B721605	B721607	B721610	B721611	B721612	B721614	B721615	B721616	B721617
	Shipment No.	8P	4P	4P	4P	4P	8P	4P	4P	8P	8P	4P	8P	8P	8P	4P	4P	8P	4P	4P	4P	4P

Results of Prescreen Testing*

Shipment No.											
	Prescreen No.	Ž,	VE	T.	AVS No.	VF	je	VE	PT	SF	W
8P	B721618	0	3	32	8503	1	0	0	0	0	1
4P	B721619	5	1	45	7401	0	0	0	0	0	0
8P	B721621	0	1	30	8324	0	0	0	0	0	0
4P	B721625	9	2	40	7402	0	0	0	0	0	0
4P	B721628	3	-	39	7403	0,1	0,0	0,2	1,1	0,0	0
4P	B721630	9	1	41	7404	0	0	0	0	0	0
12P	B721643	15	12	17	9172	1	1	2	0	1	0
4P	B721645	22	1	24	8338	1	0	0	0	0	0
1P	B721690	16	-	06	6241	0	0	0	0	0	NR
IP	B721693	61	~	62	6242	0	0	0	0	0	NR
IP	B721702	54	-	69	6243	0,1	0,0	0,0	3,0	0,0	NR
1P	B721714	52		70	6244	0	0	0	0	0	NR.
1P	B721728	74	3	78	6245	0	0	0	1	0	X
1P	B721729	22	1	49	6246	0,1	0,0	1,1	1,0	0,1	N.
10P	B721746	11	9	0	9173	2	0	2	2	2	0
1P	B721749	5	19	100	6584	0,1	0,0	1,0	1	0,0	NR
11P	B721754	51	1	77	6247	0	0	0	1	0	R
10P	B721777	0	31	0	9174	0	0	1	0	0	•
10P	B721778	16	9	0	8379	0	0	1	0	0	•
10P	B721781	12	5	46	8380	1	0	0	1	0	0
10P	B721786	0	81	14	9175	0	0	7	0	•	•

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Shipment No.	Prescreen No.	YF	VE	P	AVS No.	YF	JE	VE	M	SF	X
3Р	B721787	10	1	91	6585	0	0	0	0	0	0
3P	B721818	23	1	29	6586	0	0	0	0	0	0
1P	B721823	82,59	1	100,100	6248	0,0	0,0	1,0	2,0	0,0	0
3Р	B721826	31	1	99	6587	0	0	0	0	0	0
3P	B721838	61	1	99	6588	1,1	0,0	0,0	1,0	0,0	0
9P	B721860	0	11	24	9176	1	0	1	0	0	1
9P	B721863	0	23	38	8516	1	0	1	1	1	1
9P	B721877	10	1	15	9177	0	0	0	0	0	0
3P	B721880	90	1	92	6889	2,0	0,0	0,0	1,0	1,0	0
9P	B721881	0	1	2	8517	1	0	1	0	0	0
9P	B721883	0	2	23	8518	0	0	0	0	0	0
3Р	B721892	12	1	50	06290	0	0	0	0,0	0	0
3Р	B721899	100	1	19	6591	1,1	0,0	0,0	1,0	1,0	0
3Р	B721905	15	99	28	6592	1,1	0,0	0,0	1,0	1,0	0
зъ	B721908	1	1	50	6593	0	0	0	0	0	0
3P	B721910	32	1	86	6594	1,0	0,0	0,0	1,0	0,0	0
3Р	B721917	0	1	76	6595	0	0	0	0	0	0
3Р	B721925	9	1	29	9659	0	0	0	0	0	0
3Р	B721953	5,99	1	38,84	6597	0	0	0	0	0	•
3P	B721958	0	1	58	6598	0	0	0	0	0	0
3P	0701070	9,									

Results of Prescreen Testing^a

W	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	•
SF	0	0	0	0,0	0	0,0	0	0	0	1,0	1,0	0,0	1,0	0,0	0	0	0	0	0	0,0	•
PT	0	0	0	2,0	0	2,0	0	0	0	1,0	1,0	0,0	2,0	1,0	0	0	1	1	0	0,0	_
VE	0	0	1	0,0	0	0,0	1	0	0	0,0	1,1	0,0	0,0	0,0	0	0	0	0	0	0,0	<
JE	0	0	0	0,0	0	0,0	0	0	0	0,0	1,0	0,0	0,0	0,0	0	0	0	0	0	0,0	c
YF	0	3	0	1,0	0	1,0	0	1	0	1,1	1,1	1,1	1,1	1,1	0	0	0	0	0	1,0	-
AVS No.	0099	8519	9178	1099	2099	6603	9179	9180	1816	6604	9099	9099	2099	8099	6099	0199	1199	6612	9182	6613	7133
М	99	34	4	79	89	87	25	55	0	7.1	99	68	57	55	7.1	50	98	63	5	51	47
VE	1	21	91	1	1	1	13	2	1	1	ı	1	-	1	1	1	1	ı	2	1	
Y.	29	7	11	1	3	32	28	26	13	23	41	36	47	30	59	3	8	10	6	52	76
Prescreen No.	B722006	B722033	B722038	B722048	B722052	B722054	B722055	B722060	B722072	B722076	B722077	B722078	B722080	B722081	B722087	B722089	B722091	B722094	B722098	B722109	D722111
Shipment No.	3P	9P	9P	3P	3Р	3Р	9P	9P	9P	3Р	3Р	3Р	3Р	3P	3P	3P	3Р	3Р	9P	3Р	30

		III COM	201110	Simple of Flactical Lating			Medica	Medulis of Fillinary Serven Testing		9	
Shipment No.	Prescreen No.	YF	VE	£	AVS No.	YF	2	VE	Ē	SF	A
9P	B722116	73	3	55	9183	2	1	0	0	1	1
9P	B722117	46	27	58	9184	2	2	0	1	2	0
3P	B722141	17	-	63	6615	0	0,0	0	0	0	0
9P	B722161	2,5	1,5	31,34	9185	1	0	0	0	1	1
3P	B722162	12	_	55	9199	1	0	0	0	0	0
3P	B722165	79	1	58	6617	2,0	2,0	0,0	0,0	0,0	0
3P	B722168	23	1	82	8199	2,0	0'0	0,0	0,0	0,0	0
9P	B722172	2	25	14	9186	0	0	1	0	0	0
9P	B722174	14	10	30	8366	0	0	0	2	0	0
9P	B722174	14	10	30	2816	0	0	0	0	0	0
9P	B722179	25	1	17	9188	1	1	0	0	1	1
3P	B722181	51	1	51	6199	1	0	0	0	0	0
3P	B722182	88	-	58	6620	2	0	0	0	0	0
3P	B722183	90	1	76	6621	1	0	0	0	0	0
9P	B722184	0	22	49	8367	1	0	1	1	1	1
9P	B722186	23	1	17	9189	1	0	0	0	0	0
9P	B722190	22	0	4	9190	0	0	0	0	0	0
9P	B722194	20	1	4	1616	0	0	0	0	0	0
9P	B722198	0	30	12	9192	1	0	1	0	0	0
9P	B722199	0	28	28	9193	1	0	1	0	0	0
9P	B722210	1	2	13	8363	0	0	0	0	0	0

Results of Prescreen Testing^a

Results of Primary Screen Testing^b

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Shipment Prescreen YF VE PT AVS YF PT RF AVS YF PT SF PT No. No. DF PT SF No.			Kesul		s of Prescreen Testing		ALL PROPERTY.	Kesuits	Results of Primary Screen Testing	Screen	esting	STATE STATE
B722215 15 1 10 9194 0 0 0 0 B722222 46 - 90 6622 0,0 0,0 0,0 1,0 B722224 46 - 79 6622 0,0 0,0 0,0 1,0 B722229 2,30 12 25,5 9195 1 0 0 0 B722230 23,1 35 72,35 6625 0,0 0,0 0 <td< th=""><th>Shipment No.</th><th>Prescreen No.</th><th>YF</th><th>VE</th><th>T</th><th>AVS No.</th><th>YF</th><th>JE</th><th>VE</th><th>Ħ</th><th>SF</th><th>W</th></td<>	Shipment No.	Prescreen No.	YF	VE	T	AVS No.	YF	JE	VE	Ħ	SF	W
B722224 46 90 6622 0,0 0,0 0,0 1,0 B722224 4 79 6623 1,0 0,0 0 1,0 B722228 49 91 6624 0 0 0 1,0 B722236 2,30 12 25,5 9195 1 0 1 0 B722230 2,31 35 72,35 6625 0,0 0 0 1,0 0 0 1,0 0 0 0 1,0 0 0 1,0 0	9P	B722215	15	1	10	9194	0	0	0	0	0	0
B72224 4 - 79 6623 1,0 0,0 0 1,0 B72228 49 - 91 6624 0 0 0 0 B72228 2,30 12 25,5 9195 1 0 1 0 B72230 2,31 35 72,35 6625 0,0 0 0 1,0 B722234 0,13 17 40,38 8368 1 0 1 0 B722234 11 23 21 9196 0	3P	B722222	46	l	90	6622	0,0	0,0	0,0	1,0	0,0	0
B722228 49 - 91 6624 0 0 0 0 B722229 2,30 12 25,5 9195 1 0 1 0 B722230 23,1 35 72,35 6625 0,0 0,0 0 1,0 B722231 0,13 17 40,38 8368 1 0 1 0 B722233 26 0	3P	B722224	4	_	79	6623	1,0	0,0	0	1,0	0,0	0
B722230 2,30 12 25,5 9195 1 0 1 0 B722230 23,1 35 72,35 6625 0,0 0,0 0 1,0 B722231 0,13 17 40,38 8368 1 0 1 2 B722233 26 0 </td <td>3P</td> <td>B722228</td> <td>49</td> <td>-</td> <td>91</td> <td>6624</td> <td>0</td> <td>0</td> <td>0</td> <td>0</td> <td>0</td> <td>0</td>	3P	B722228	49	-	91	6624	0	0	0	0	0	0
B722230 23,1 35 72,35 6625 0,0 0,0 0,0 1,0 B722231 0,13 17 40,38 8368 1 0 1 2 B722233 26 0 0 9196 0 0 0 0 B722234 11 23 21 9197 0 0 1 0 B722236 19 0 6 9198 0 0 0 0 B722239 18,9 2 59,32 6626 0	9P	B722229	2,30	12	25,5	9195	1	0	1	0	0	0
B722231 0,13 17 40,38 8368 1 0 1 2 B722233 26 0 0 9196 0	3P	B722230	23,1	35	72,35	6625	0,0	0,0	0	1,0	0,0	0
B722234 26 0 0 9196 0 0 0 0 B722234 11 23 21 9197 0 0 1 0 B722236 19 0 6 9198 0 0 0 0 B722230 78,9 2 59,32 6626 0 0 0 0 0 B722240 16 4 29 8364 3,1 2,0 0,0 1,1 0 <td< td=""><td>9P</td><td>B722231</td><td>0,13</td><td>17</td><td>40,38</td><td>8368</td><td>1</td><td>0</td><td>1</td><td>2</td><td>1</td><td>1</td></td<>	9P	B722231	0,13	17	40,38	8368	1	0	1	2	1	1
B722234 11 23 21 9197 0 0 1 0 B722236 19 0 6 9198 0 0 0 0 B722239 78,9 2 59,32 6626 0 0 0 0 B722240 16 4 29 8364 3,1 2,0 0,0 0 0 0 B722241 100 - 78 6627 0 0 0 0 0 0 0 B722242 6,47 2 45,6 9199 1 2 0 </td <td>9P</td> <td>B722233</td> <td>26</td> <td>0</td> <td>0</td> <td>9616</td> <td>0</td> <td>0</td> <td>0</td> <td>0</td> <td>0</td> <td>0</td>	9P	B722233	26	0	0	9616	0	0	0	0	0	0
B722236 19 0 6 9198 0 0 0 0 B722239 78,9 2 59,32 6626 0 0 0 0 0 B722240 16 4 29 8364 3,1 2,0 0,0 0 0 0 0 B722241 100 - 78 6627 0	9P	B722234	11	23	21	9197	0	0	1	0	1	0
B722240 78,9 2 59,32 6626 0 0 0 0 0 B722240 16 4 29 8364 3,1 2,0 0,0 1,1 B722241 100 - 78 6627 0 0 0 0 B722242 6,47 2 45,6 9199 1 2 0 0 0 B722245 0,0 25 17,9 9200 0 1 0 0 0 0 B722246 63,10 4 56,36 6628 2 0 0 0 0 B722247 61,0 2 56,3 6629 1 0 0 0 0 0 B722248 0,64 5 29,3 9202 1 2 0 0 0 0 B72255 0 1 1 1 1 1 0 0 0	9P	B722236	19	0	9	9198	0	0	0	0	0	0
B722240 16 4 29 8364 3,1 2,0 0,0 1,1 B722241 100 - 78 6627 0 0 0 0 B722242 6,47 2 45,6 9199 1 2 0 0 B722245 6,47 25 17,9 9200 0 1 0 0 B722245 6,14 0 17,0 9201 0 0 0 0 0 B722247 61,0 2 56,36 6629 1 0 0 0 0 0 B722247 61,0 2 56,2 6629 1 2 0 0 0 0 B722248 0,64 5 29,3 9202 1 2 0 0 0 0 B722255 0 26 0 9204 1 1 1 1 0 0	3P	B722239	78,9	2	59,32	9799	0	0	0	0	0	0
B722241 100 - 78 6627 0 <	9 P	B722240	16	4	29	8364	3,1	2,0	0,0	1,1	1,0	0
B722242 6,47 2 45,6 9199 1 2 0 0 B722244 0,0 25 17,9 9200 0 1 0 <t< td=""><td>3P</td><td>B722241</td><td>100</td><td>1</td><td>78</td><td>6627</td><td>0</td><td>0</td><td>0</td><td>0</td><td>0</td><td>0</td></t<>	3P	B722241	100	1	78	6627	0	0	0	0	0	0
B722244 0,0 25 17,9 9200 0 1 0	9 P	B722242	6,47	7	45,6	6616	1	2	0	0	0	0
B722245 0,14 0 17,0 9201 0 0 0 0 B722246 63,10 4 56,36 6628 2 0 0 0 0 B722247 61,0 2 56,2 6629 1 0 0 0 0 B722248 0,64 5 29,3 9202 1 2 0 0 0 B722255 0 26 0 9203 1 1 1 0 0 B722256 0 13 0 9204 1 1 1 0 0	9P	B722244	0,0	25	17,9	9200	0	1	0	0	0	0
B722246 63,10 4 56,36 6628 2 0 0 0 B722247 61,0 2 56,2 6629 1 0 0 0 B722248 0,64 5 29,3 9202 1 2 0 0 B722255 0 26 0 9203 1 1 0 0 B722256 0 13 0 9204 1 1 1 0	9P	B722245	0,14	0	17,0	9201	0	0	0	0	0	0
B722247 61,0 2 56,2 6629 1 0 0 0 B722248 0,64 5 29,3 9202 1 2 0 0 B722255 0 26 0 9204 1 1 0 0	3P	B722246	63,10	4	56,36	6628	2	0	0	0	0	0
B722248 0,64 5 29,3 9202 1 2 0 0 B722255 0 13 0 9204 1 1 1 0	3P	B722247	0,19	2	56,2	6629	1	0	0	0	0	-
B722255 0 26 0 9203 1 1 0 0 B722256 0 13 0 9204 1 1 1 0	9P	B722248	0,64	5	29,3	9202	1	2	0	0	0	0
B722256 0 13 0 9204 1 1 1	9P	B722255	0	26	0	9203	1	1	0	0	1	0
	99	B722256	0	13	0	9204	-	1	1	0	-	•

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	The state of the s		Summer in the second	Control of the last		NESOIL	Reulis of Friniary Screen Leaning	ושחה /	Sime	ALTER SALES
Prescreen No.	, VR	VE	PT	AVS No.	YF	JE	VE	Id	SF	W
B722278	13	2	0	8365	0	0	0	0	0	0
B722279	13	1	09	8238	0	0	0	1	0	1
B722280	5	0	63	8239	0	0	0	1	0	2,2
B722504	S	32	80	9205	1	0	1	0	0	0
B722506	4	22	1	8381	0	0	0	0	0	0
B722508	6	20	41	8382	1	0	1	1	0	0
B722510	0	37	ĸ	8383	1	0	1	0	0	0
B722518	84	-	0	7109	0,1	0'0	0,0	0'0	0,0	0
B722525	0	-	79	7110	0,0	0,0	1,0	1,0	0,2	3,2
B722559	28	14	99	8240	2,2	0,0	1,2	2,2	1,2	1
B722566	29	9	14	8270	1,1	0,0	2,2	2,1	0,1	1
B722591	15	3	51	8241	1,1	0,0	2,1	1,1	1,0	1
B722607	0	2	54	8242	0	0	1	0	0	0
B722614	1	4	90	8243	I	0	0	0	0	0
B722628	1	3	19	8244	1	0	1	0	0	0
B722632	0	1	LS.	8245	0	0	0	0	0	0
B722634	4	1	65	8246	0	0	0	0	0	0
B722635	20	1	12	8512	1	0	1	0	0	0
B722666	0	1	28	8247	1	0	1	0	0	0
B722668	1	5	42	8248	1,1	0,0	1,1	0,1	1,0	1
B722689	2	15	<i>L</i> 9	8228	1,2	0,0	0,1	2,2	1,2	0

Results of Prescreen Testing

3	0,0	0	0	0	1	1	0	0	1	0	0	0	0	0	1	0	0	0	0	0	0
SF	0,0	2,1	2,2	1,0	0,1	0	-	0	0	0	3,2	0	0	1,0	1,0	0,0	0	0	0	0	0,0
PT SE	0,2	2,1	2,2	2,0	2,1	0	1	0	0	0	2,2	0	0	2,2	0,0	0,0	0	0	0	0	1,0
JE VE PT	0,0	0,0	1,1	0,0	1,1	1	1	0	0	0	0,0	0	0	0,0	0,0	0,0	0	0	0	1	1,1
63																					
JE	0,0	0,0	0,0	0,0	0,0	0	0	0	0	0	0,0	0	0	0,0	1,1	0,0	0	0	0	0	0,0
YE	0,0	0,0	2,0	0,0	1,0	0	1	0	0	0	2,2	0	0	0,1	1,1	0,0	0	0	•	0	2,1
AVS No.	8339	8250	8251	7405	8229	8252	8253	7111	7112	7113	8271	7114	7115	7406	7407	8340	7312	7313	8413	9208	8373
H.	0	89	99	0	13	33	49	57	19	20	73	100	55	19,28	13,30	22,36	49,54	71,0	25	25	2
VE		10	11	1	17	8	6	1	1	1	1	1	1	-	1	1	1	1	18	18	7
YF	25	4	1	61	0	0	0	1	5	3	3	100	7	11,18	9,18	0,20	0,0	24,9	8	8	36
Prescreen No.	B722733	B722743	B722745	B722752	B722757	B722778	B722785	B722805	B722808	B722811	B722823	B722824	B722849	B722854	B722858	B722866	B722867	B722871	B722872	B722872	B722873
Shipment No.	7P	<i>T</i> P	<i>T</i> P	3P	<i>TP</i>	7P	7P	3P	3P	3P	7P	3P	3P	3P	3P	7P	3P	3P	10P	10P	10P

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		Results		of Prescreen Testing*			Results	of Primar	Results of Primary Screen Testing ^b	esting	
Shipment No.	Prescreen No.	YF	VE	M	AVS No.	YF	æ	VE	Ħ	SF	A
10P	B722873	36	7	64	6076	1	0	1	0	0	0
3P	B722874	3,2	-	76,30	7314	0,0	0,0	0,0	1,0	0,0	0
10P	B722876	15	0	2	8388	0	0	0	0	0	0
10P	B722876	15	0	2	9210	0	0	0	0	0	0
10P	B722883	9,5,2	15,2	16,55	7315	0	0	0	1	0	0
3P	B722883	9,5,2	15,2	16,55	7315	0	0	0	1	0	0,0
10P	B722884	12	0	10	8389	0	0	0	0	0	0
10P	B722884	12	0	10	9212	1	0	0	0	0	0
10P	B722886	8,0,0	3	21,61,40	7316	0	0	0	1	0	0,0
3P	B722886	8,0,0	28	21,61,40	7316	0	0	0	1	0	0,0
3P	B722889	4,3,8	28	19,51,17	7317	0	0	0	0	0	0,0
10P	B722889	4,3,8	28	19,51,17	7317	0	0	0	0	0	0,0
10P	B722890	13	31	61	8390	0	0	0	0	0	0
10P	B722890	13	31	61	9215	0	0	0	0	0	0
10P	B722899	0	26	34	9216	0	0	1	0	0	0
10P	B722902	8	5	22	8374	3,2	2,1	2,2	2,1	3,2,2	0
10P	B722902	8	5	22	9217	2	1	1	1	8	0
3P	B722904	6,13	_	33,52	7318	0	0	0	1	0	0
10P	B722905	80	17	3	9218	0	0	0	0	0	0
10P	B722908	0	19	0	9219	1	0	1	0	0	0
10P	B722909	22	80	4	8391	0,0,0	0,0,0,0	0,1,0	0,0,1	0,0,0	0

Results of Prescreen Testing

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Shipment No.	Prescreen No.	YF	VE	73	AVS No.	YF	ær	VE	М	SF	W
10P	B722911	0	23	5	8220	1	0	0	0	0	0
10P	B722914	1	39	0	9221	1	0	7	0	0	0
10P	B722915	4	29	37	8375	0,0	0,0	0,1	2,1	0,0,0	0
10P	B722915	4	29	37	9222	1	0	1	0	0	0
10P	B722917	3	25	4	9223	1	0	0	0	0	0
10P	B722921	25	14	25	8376	1,1	1,0	0,1	2,2	2,1	1
10P	B722921	25	14	25	8392	NT	INT	NT	IN	IN	NT
10P	B722922	4	12	9	8337	1	1	1	0	0	2
10P	B722922	4	12	9	9224	0	0	0	0	0	0
10P	B722926	9	27	40	9225	1	. 2	1	7	0	0
10P	B722929	1	15	2	8414	0	0	1	1	0	0
10P	B722931	12	20	*	9226	0	0	1	1	0	0
10P	B722932	7	30	15	9227	0	2	1	1	0	0
10P	B722933	20	8	3	8364	3,1	2,0	0'0	1,1	1,0	0
10P	B722935	4	21	19	9228	0	0	1	1	0	0
10P	B722936	9	42	62	9229	1	0	1	7	0	0
10P	B722937	21	10	17	8369	1	0	1	0	0	0
10P	B722938	9	30	13	9230	0	0	1	0	0	0
10P	B722942	13	33	11	8386	0	0	1	0	0	0
10P	B722944	9	29	9	9231	0	0	0	0	0	0
10P	B722950	9	27	10	9232	1	0	1	0	0	0

Bull
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Results

¥	1	0	0	0	0	0,0	0	0	0	0	M	FN.	0	0	Į.	0	0	0	ĮN.	0	NT
SF	3,2	3	0	1	0	0,0	0,0	0,0	0	0,0	0	0,0	0,0	0,0	1,1	0,0	1,0	1,0	0,0	1	0
PT	1,1	0	1	2	0	0,0	1,1	1,2	0	1,2	0	0,2	0,0	0,0	0,2	2,0	0,1	2,0	0,1	0	0
VE	1,0	0	0	0	2	0,0	1,1	1,2	0	1,0	0	0,0	0,0	0,0	1,0	0,0	0,0	0,0	0,0	0	0
JE	0,0	0	0	2	0	0,0	0,0	0,0	0	2,2	0	2,3	0,0	0,0	0,0	0,0	0,0	0,0	1,1	1	0
YF	2,0	0	0	1	0	0,0	2,1	0,0	0	2,2	0	0,2	0,0	0,0	1,0	0,0	0,0	0,0	2,2	1	0
AVS No.	8370	9233	9234	9235	9236	8341	8371	7116	21112	8372	7443	7445	9618	8197	7439	7408	7409	7410	7438	8342	7446
M	21	21	4	3	0	6	27	99	52	29	0	54	7	00	10	0	1	14	6	10	3
VE	32	32	20	30	20		39		1	19	63	28	49	22	84	•			5		53
YF	10	10	8	8	4	27	7	6	15	12	0	43	0	5	2	14	13	7	72	27	7
Prescreen No.	B722965	B722965	B722976	B722987	B722992	B722997	B723035	B723037	B723044	B723047	B723061	B723062	B723064	B723089	B723096	B723106	B723107	B723110	B723123	B723126	B723136
Shipment No.	10P	10P	10P	10P	12P	3P	10P	3P	3P	10P	SP	SP	SP	SP	SP	3P	3P	3P	SP	3P	SP

esults of Prescreen Testing

	Total State of the	Result	s of Prescr	is of Prescreen Testing	Section 1		Results	Results of Primary Screen Testing	y Screen T	esting"	
Shipment No.	Prescreen No.	YF	VE	m	AVS No.	YF	JE	VE	PT	SF	W
SP	B723139	67	8	5	8262	2,1	0,0	0,2	2,1	1,0	0
SP	B723141	100	69	3	7440	0	0	0	0	0	M
SP	B723143	2	9	0	7442	0	0	0	0	0	K
SP	B723148	3	73	0	7441	0	0	0	0	0	¥
SP	B723159	0	49	0	8198	0	0	1,0	0,1	0,1	1
5P	B723160	0	28	0	8199	0	0	0,1	1,0	1,0	0
SP	B723169	48	14	27	9238	1	1	0	2	0	0
SP	B723172	4	1	26	9239	0	0	0	0	0	0
SP	B723199	35	91	14	8263	2,1	2,2	1,0	2,2	0,1	0
SP	B723201	28	51	0	8264	1	0	0	0	0	0
5P	B723203	0	09	9	7444	0,0	0,0	1,2	0,0	0,0	F
SP	B723223	0	45	4	8332	2,1	0,0	1,1	2,1	2,1	1
5P	B723230	21	4	2	8265	1	0	0	0	0	0
5P	B723233	0	38	3	8200	0	0	1,1	0,0	0,0	1
5P	B723240	0	59	0	8333	1	0	1	0	0	1
SP	B723245	3	9	48	8232	0	0	0	0	0	0
5P	B723247	0	71	34	8201	0	0	1,1	0,0	1,0	0
Ф.	B723250	1	64	1	8212	0,2	0,0	2,2	1,2	1,1	1
Ф.	B723256	0	1	27	8213	0	0	1	1	0	0
Ф.	B723261	0		29	8214	0	0	1	0	0	0
GP	B723262	12	1	77	8215	0	0	1	0	0	0

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	X	0	0	0	1	0	1	0	0	0	1	0	0	0		0	0 1	0 1 0	0 1 0	0 1 0 0 1	0 1 0 0 1 0	0 1 0 0 1 0 0
Sting	SF	1	0,1	1,0	1,0	0,0	1,0	0	0,1	1,1	0	0,0	0,2	0,0		0	0 -	0,0	0,0 0,1	0 0,0 0,1 0	0 0,0 0,0	0 0,0 0,0 0,0
Newton of Lineary Server Testing	K	1	0,0	1,0	2,1	1,1	2,1	1	1,1	1,1	1	2,1	1,1	1,1			1 1	0,0	0,0	0,0	0,0 0,0 0,0	0,0 0,0 0,0
	VE	0	0,0	0,0	1,1	0,0	0,1	1	2,3	1,1	1	1,1	1,1	0,0	1		1	0,0	0,0	0,0	0,0 1,1 0 1,0	1,1 0,0 0 1,0 0,0
-	JE	0	0	0	0,0	1,0	0,0	0	2,2	0,0	0	0,0	0,0	0,0	0		0	0	0,0	0 0,0	0 000	0 0 0 0 0
	£	1	1	1	1,1	2,1	0,1	0	1,1	1,1	0	1,1	2,1	2,2	0		0	0 0	0 0 1	0 0 1	0 0 1 1 0	0 0 1 1 0 0
	AVS No.	8267	8216	8217	8218	8268	8219	8220	8221	8222	8223	8224	8269	8228	8226		8227	8227	8227 8210 8211	8227 8210 8211 8266	8227 8210 8211 8266 8202	8227 8210 8211 8266 8202 8203
0	E.	36	51	49	57	42	17	39	39	59	73	39	29	7	14	•	0	33	33	33 42 32	33 42 32 37	33 42 32 37 25
	VE	1	1	1	1	1	1	1	1	1	-	1	1	28	74	9	3	78	78	57 57 10.	38 10.01	78 10. 10. 27
	YF	19	11	49	6	45	3	23	21	28	3	35	30	48	0	7		0	38	38	38 31 1	0 38 11 1 1
	Prescreen No.	B723266	B723268	B723275	B723278	B723279	B723285	B723286	B723289	B723290	B723297	B723298	B723300	B723306	B723315	B723318		B723322	B723322 B723364	B723322 B723364 B723398	B723322 B723364 B723398 B723400	B723322 B723364 B723398 B723400 B723401
	Shipment No.	6P	6P	6P	6P	9P	GP.	6P	6P	6P	6P	6P	6P	6P	6P	6P		6P	6P 6P	6P 6P	6P 6P 6P	6P 6P

Testing
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	W	0	0	0	1	0	0	0	0	0	0	0	0	0	. 0	0	0	0	0	0	0	C
Summary of a summary of the summary	SF	0,0	0,0	0,0	1,0	0	0	0	0	0	1	0	0	2,2	0	0	3	1	0	0	0	•
	PT	1,1	1,0	0,0	1,1	0	0	1	0	0	0	0	1	3,2	2	0	2	0	0	0	0	_
	VE	0,1	0,0	0,0	1,1	0	0	0	0	0	0	0	0	1,1	1	0	1	-	0	0	1	
	JE	1	1	0	0	0	0	0	0	0	0	0	0	0,0	0	0	2	0	0	0	0	•
	YF	1	1	0	0	0	1	0	0	0	1	0	0	2,1	0	2	2	0	1	1	1	•
	AVS No.	8205	8206	8207	8208	9240	9241	9242	9243	8393	8394	9244	9245	8378	9246	9247	9248	9249	9250	9251	8403	0262
	Ŗ	35	46	35	26	5	2	2	4	16	24	0	2	99	18	0	11	8	3	0	4	P
	VE	43	22	36	75	14	24	22	27	21	6	18	30	99	45	. 5	4	2	2	5	26	12
	YF	22	53	22	0	5	Û	0	1	51	42	0	0	15	9	26	50	19	55	15	0	0
	Prescreen No.	B723410	B723412	B723413	B723414	B723420	B723424	B723425	B723426	B723427	B723428	B723429	B723430	B723431	B723432	B723435	B723436	B723438	B723441	B723444	B723448	B723440
	Shipment No.	6P	GP	GP	GP	12P	10P	100														

2 0 0 1 1 0 1 1			0 0 0 0 0,0 0,1	0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	0 1 0 0,0 0,0 0 0,0 0,0 0,0 0,0 0,0 0,0	0 1 0 0,0 0,0 0 0,0 0,0 0,0 0,0 0,0 0,0	0 1 0 0,0 0,0 0,0 0,0 0,0 0,0 0,0	0 1 0 0,0 0,0 0 0,0 0,0 0,0 0,0 0,0 0,0	0 1 0 0,0 0,0 0 0,0 0 0,0 0 0 0 0	0 1 0 0,0 0,0 0 0 0 0 0 0 0 0 0 0 0 0 0	0 1 0 0,0 0,0 0 0 0 0 0 0 0 0 0 0				
	0 0,0	0 1 1 0,0	0 1 1 0,0 1,1 0	0 1 1 0,0 1 1	0 1 1,1 0,0 1 0	0 1 1 0,0 1 0 0,0 0,0 0,0	0 1 1 0,0 0 0,0 0 2	0 1 1 0,0 1,1 0,0 0,0 2 0,0 0,0 0,0	0 1 1 0,0 1,1 0,0 0,0 2 0,0 2	0 1 1 0,0 1,1 0,0 0 2 0,0 2 0 0,0 0 0 0 0 0 0 0 0 0 0	0 1 1,1 0,0 0,0 0 0,0 0 0 0 0 0 0 0 0 0	0 1 1,1 0,0 0,0 0 0 0 0 0 0 0 0 0 0 0	0 1 1 0,0 1,1 0,0 0 0 0 0 0 0 0 0 0 0 0	0 1 1 0,0 1,1 1,1 0 0,0 2 0,0 2 0 0 0 0 1	0 1 1 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	
0 1																
0 -	0,0	0,0	0,	0 0,0 0,0	0 0,0 0,0 0	0 0,0 0,0 0,0	0 0,0 0,0 0	0 1 0,0 0 0 0,0 0 0,0	0 1 0,0 0 0 0,0 0	0 1 0,0 0,0 0 0,0 0	0 1 0,0 0,0 0 0,0 0	0 1 0,0 0,0 0 0,0 0 0	0 1 0,0 0,0 0 0,0 0 0	0 1 0,0 0,0 0 0,0 0 0 0 0	0 1 0,0 0,0 0 0,0 0 0 0 0 0 1	0 1 0,0 0,0 0 0,0 0 0 0 0 0 1 1
0	0 2,1	2,1	2,1	0 2,1 1,1,1 1,1	0 1,1,1 1,1 0 0	0 1,1,1 0 0 1,1	0 2,1 1,1,1 1,1,1 0 0 0 0 0	0 1,1,1 1,1,1 0 0 0,1 0,1	0 1,1,1 1,1,1 0 0 0,1 0 0	0 1,1,1 1,1,1 0 0 0 1,0 0 0 1,0	0 1,1,1 1,1,1 0 0 0,1,0 0 0 0 0 0	0 1,1,1 1,1 0 0 1,1 0 0 0 0 0 0 0 0 0 0	0 1,1,1 1,1 0 0 1,1 0 0 0 0 0 0 0 0 0 0	0 1,1,1,1 0 0 1,1,0 0 0 0 0 0 0	0 1,1,1,1 0 0 1,1 0 0 0 0 0 0 0	0 1,1,1,1 0 0 1,0 0 0 0 0 0 0
9255	2 2	9 2 2					8 8 7 9 7 8 8	8 8 9 7 9 7 8 8 6	8 8 9 7 9 7 8 8 8 6 6	8 8 8 6 6 0		0 1 0 0 8 8 8 7 9 7 9 8 8		1 2 0 1 0 8 8 8 7 6 7 6 2 8	2 1 2 0 1 0 8 8 8 4 6 7 6 2 8	
	53	8 8	339 839 839	8395 8396 8397 9256	8398 8394 8397 8397 8397	8396 8397 8257 8398	83 83 83 83 83 83 83		83 83 82 83 83 83 83 83 83 83 83 83 83 83 83 83	83 83 83 83 83 83 83 83 83 83 83 83 83 8	83 83 83 83 83 83 84 84 84 85 85 85 85 85 85 85 85 85 85 85 85 85	83 83 83 83 83 83 83 83 83 83 83 83 83 8	839 839 839 839 839 839 839 839 839 839	25 26 26 27 28 28 28 28 28 28 28	25 25 26 27 28 27 28 28 28 28 28	8396 8396 840 840 840 840
33	32	39	39	39 35	39 35 19	35 35 11	35 39 11 11 12 62	35 39 35 11 11 22 22	35 35 11 11 12 22 14	35 35 11 11 14 14	35 35 35 11 11 14 14	35 35 36 11 11 14 14 15	22 35 11 11 12 12 14 14	35 35 36 37 37 38 39 39 30 11 11 12 13 14 14 16 17 18 18 18 19 19 19 19 19 19 19 19 19 19 19 19 19	35 39 37 37 38 39 39 30 31 11 11 11 12 30 30 30 30 30 30 30 30 30 30 30 30 30	35 39 35 39 35 39 39 39 39 39 39 39 39 39 39 39 39 39
24 24 32 83		39	65 2	65 39 6 35	65 2 6 35 7 19	65 2 6 35 37 19 24 11	65 2 6 35 37 19 24 11 4 62	65 2 6 35 6 35 37 19 4 62 18 22	65 2 6 35 6 35 37 19 4 62 18 22 14	65 2 6 35 6 35 37 19 24 11 4 62 18 22 25 14 23 2	65 2 65 2 6 35 37 19 24 11 4 62 18 22 25 14 23 2 24 1	46 39 65 2 6 35 24 11 4 62 18 22 23 2 24 1 23 2 24 1 3 15	65 2 65 2 6 35 24 11 4 62 18 22 23 2 24 1 23 2 24 1 23 14 23 14 23 14 23 14	46 39 65 2 6 35 24 11 4 62 18 22 23 2 24 1 23 2 24 1 3 15 23 14 23 14 7 16	46 39 65 2 6 35 24 11 4 62 18 22 23 2 24 1 23 14 23 14 7 16 4 20	46 39 65 2 6 35 24 11 4 62 18 22 25 14 24 1 24 1 23 2 24 1 4 20 4 15
	24 32	10 46 39	10 46 39 0 65 2	24 24 32 10 46 39 0 65 2 2 6 35	24 24 32 10 46 39 0 65 2 2 6 35 7 37 19	24 24 32 10 46 39 0 65 2 2 6 35 7 37 19 6 24 11	24 24 32 10 46 39 0 65 2 2 6 35 7 37 19 6 24 11 4 4 62	24 24 32 10 46 39 0 65 2 2 6 35 7 37 19 6 24 11 4 4 62 2 18 22	24 24 32 10 46 39 0 65 2 2 6 35 7 37 19 6 24 11 4 4 62 2 18 22 5 25 14	24 24 32 10 46 39 0 65 2 2 6 35 7 37 19 6 24 11 4 4 62 2 18 22 5 25 14 4 23 2	24 24 32 10 46 39 0 65 2 2 6 35 6 24 11 6 24 11 4 4 62 2 18 22 5 25 14 4 23 2 3 24 1	24 24 32 10 46 39 0 65 2 2 6 35 6 24 11 6 24 11 2 18 22 5 25 14 4 23 2 3 24 1 8 3 15	24 24 32 10 46 39 0 65 2 2 6 35 6 24 11 6 24 11 6 24 11 5 18 22 5 25 14 4 23 2 3 24 1 8 3 15 3 23 14 3 23 14	24 24 32 10 46 39 0 65 2 2 6 35 4 4 6 24 4 4 62 4 4 62 5 25 14 4 23 2 3 24 1 8 3 14 8 3 14 6 7 16	24 24 32 10 46 39 0 65 2 2 6 35 4 4 62 4 4 62 4 4 62 5 25 14 4 23 2 3 24 1 8 3 15 8 3 14 6 7 16 19 4 20	24 24 32 10 46 39 0 65 2 2 6 35 4 4 6 24 4 4 62 4 4 62 4 4 6 2 4 4 6 14 4 23 2 14 8 3 14 1 6 7 16 16 19 4 20 44 4 15

Results of Prescreen Testinga

W	0	0	1	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	•	0
SF	1	0	0	0	1	0	0	0,0	0,0	0	0	0,0	0	0	1,0	0	0	0	0,0	1,0	0,0
М	1	1	0	0	2	0	0	0	0	0	0	0.	O	0	0,0	0	0	0	1,0	0,0	0,0
VE	1	0	2	2	1	0	0	0,0	0,0	0	0	0'0	0	0	0,0	0	0	0	0,0	0,0	0,0
JE	0	0	0	0	0	0	0	0,0	0,0	0	0	0,0	0	0	0,0	0	0	0	0,0	0,0	0,0
YF	0	0	2	0	0	0	0	2,0	1,0	0	0	1,0	0	0	0,1	0	0	0	1,0	1,0	0,1
AVS No.	8265	9566	9267	9268	9269	0630	6631	6632	6633	6634	6635	9636	6637	6638	6639	6640	6641	6642	6643	6644	6645
PT	24	19	16	7	48	100	99	70	70	53	65	09	51	99	20	50	74	72	75	84	80
VE	72		16	79	62	90	1	1	1	-	-	,	1	Į:	•			1		-	1
Y.	4	0	14	4	5	2	61	15	8	9	20	6	ø	4	3	1	5	111	1	0	s
Prescreen No.	B723487	B723493	B723801	B723802	B723807	B724373	B724379	B724382	B724384	B724385	B724387	B724394	B724396	B724405	B724406	B724411	B724413	B724415	B724416	B724417	B724418
Shipment No.	401	10P	10P	10P	10P	11	dI	dI	ПР	IP .	1P	di	1.6	1.6	IP	1P	1P	IP	IP	1P	11

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		Results	1.6	of Prescreen Testing ^a			Results	Results of Primary Screen Testing ^b	y Screen T	esting ^b		
nipment No.	Prescreen No.	YF	VE	Z	AVS No.	ŸF	JE	VE	¥	SF	W	
11	B724420	0	1	0	6646	1,0	0,0	0,0	1,0	0,0	0	
11P	B724423	25	1	77	6647	0	0	0	0	0	0	
1P	B724433	7	1	77	6648	1,0	0,0	0,0	0,0	0,0	0	
11P	B724434	6	-	19	6649	0	4	0	0	0	0	
11P	B724436	28	-	55	6650	0	0	0	0	0	0	
1P	B724439	0	-	56	6651	1,0	0,0	0,0	0,0	0,0	0	
1P	B724442	59	1	64	6652	0	0	0	0	0	0	
11	B724447	10	-	64	6653	1,0	0,0	0,0	1,0	1,0	0	
1P	B724453	7	1	67	6654	1,0	0,0	0,0	0,0	1,0	0	
1P	B724455	12	-	78	6655	1,0	0,0	0,1	0,0	0,0	0	
11	B724456	30	1	96	9599	0,0	0,0	0,0	1,1	1,0	0	
11P	B724457	17	-	86	6657	1,0	0,0	0,0	1,1	0,0	0	
1P	B724458	12	-	88	8599	0	0	0	0	0	0	
1P	B724466	75	1	73	6599	0	0	2,0	0	1	0	
11P	B724468	50	-	36	0999	0,0	0,0	1,1	0,1	0,0	1	
119	B724508	6	1	65	1999	0	0	0	0	0	0	
119	B724509	30	-	52	6662	0	0	0	0	0	0	
11	B724512	111	1	62	6663	0	0	0	1	0	0	
11P	B724517	23	-	57	6664	1	0	0	0	0	0	
11	B724519	19	-	76	9999	0	0	0	0	0	0	
11	B724521	4	-	97	9999	0	0	0	0	0	0	

Results of Prescreen Testinga

Results of Primary Screen Testing^b

Shipment No.	Prescreen No.	YF	VE	Z.	AVS No.	YF	æ	VE	Pľ	SF	A
1P	B724525	20	1	19	1999	0	0	0	1	0	0
1P	B724526	37	1	99	8999	1	0	0	0	0	0
1P	B724527	0	1	92	6999	0	0	0	0	0	0
1P	B724530	09	1	15	0299	0	0	0	0	0	0
1P	B724535	12	1	LS	1299	0	0	0	0	0	0
1P	B724544	16	-	83	6672	0	0	0	0	0	0
1P	B724546	14	1	42	6249	0,0	0,0	0,0	2,1	0,0	NR
1P	B724549	0	-	05	6673	0	0	0	0	0	0
11	B724553	44	-	<i>SL</i>	6674	0	0	0	0	0	0
1P	B724558	1	1	65	6675	1,1	1,0	1,1	1,0	2,0	1
1P	B724566	25	-	85	9299	1,1	0,0	0,0	0,1	0,0	0
1P	B724586	0	ı	92	<i>CE 11</i>	0	0	0	1	0	0
1P	B724590	2	1	63	8299	0	0	0	0	0	0
1P	B724592	81	1	95	6299	1	1	0	0	0	0
11P	B724596	11	1	51	0899	0	0	0	1	0	F
1P	B724607	0	1	70	1899	1	0	0	1	0	0
11	B724610	92	1	21	6682	0	0	0	1	0	0
. 1P	B724618	88	1	24	6683	2	1	0	0	0	0
11P	B724627	18	1	89	6684	0	0	0	0	0	0
1P	B724633	3	-	54	6685	0	0	0	0	0	0
IP	B724642	5	-	79	9899	0	0	0	0	0	0

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	M	0	0	0	0	0	0	0	NR	NR	NR	NR	N.	NR.	N. N.	NR	NR.	NR	0	0	0	R.
resting ^b	SF	0,0	0	0	0	0	0	0	0,0	0,0	0	0,0	0	0,0	0,0	0	0,0	0	0	0	0	0
y Screen	K	0,0	0	0	0	0	0	0	1,1	0,1	1	0,0	0	1,0	1,0	0	1,0	0	0	0	0	0
Results of Primary Screen Testing ^b	VE	0,0	0	0	0	0	0	0	0,0	0,0	0	0,0	0	0,0	0,0	0	0,0	0	0	0	0	0
Results	Æ	0,0	0	0	0	0	0	0	0,0	0,0	0	0,0	0	0,0	1,1	0	0,0	0	0	0	0	0
	YF	0,0	0	0	0	0	0	0	1,1	0,0	0	0,0	1	0,0	2,1	0	1,1	0	0	0	0	0
	AVS No.	2899	8899	6899	0699	1699	7699	6699	6249	6251	6252	6253	6254	6255	6256	6257	6258	6229	6694	\$699	9699	6260
- 11			_			_				_					_							
een Testing	¥	61	64	63	89	09	72	55	09	72	75	33	79	19	19	70	62	83	54	92	11	41
s of Prescreen Testing ^a	VE PT	- 61	- 64	- 63	- 68	09 -	- 72	- 55	09 -	_ 72	- 75	- 33	- 79	- 61	- 61	- 70	- 62	- 83	- 54	92 -	- 71	- 41
Results of Prescreen Testing																						
Results of Prescreen Testing	VE	1	1	-	1	1	1	-	-	-	1	1	1	1	1	-	1	1	1	-	1	1

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8	1	MR	NR	NR	NR	0	NR	0	0	NR	NT	NR	X	Z	0	NR.	Z.	R	NR.	0	NR
SF	0,0	0,0	0	0	0	0	0,0	0,0	0	0	0	0	0.0	0	0,0	0	0	0	0	0	0,0
ld	0,0	0	0	0	0	0	0,0	0,0	0	0	0	0	0,0	0	6,0	0	1	0	0	0	0,0
VE	1,0	0,1	0	0	0	0	0,0	0,1	0	0	0	0	0,0	0	0,0	0	0	0	0	0	0,0
æ	0,0	0,0	0	0	0	0	0,0	0,0	0	0	0	0	1,0	0	0,0	0	0	0	0	0	0,0
K.	1,0	0,0,0	1	1	0	0	1,1	0,0	0	0	0	0	1,1	1	1,0	0	0	0	0	0	1,1
AVS No.	L699	1979	6262	6263	6264	8699	6265	6699	0029	9979	1029	6267	6268	6269	6702	6270	6271	6272	6273	6703	6274
Ł	51	85	79	82	63	1	84	79	0.0	52	33	5	06	74	61	32	56	36	55	69	61
VE	1	1	1	1	1	1	1	1	1	1	•	1	:	1	1	1	1	1	1	1	1
YF	0	47	7	28	4	58	13	17	8	0	55	3	24	38	4	71	26	09	4	10	1
Prescreen No.	B724732	B724740	B724762	B724764	B724769	B724772	B724781	B724783	B724785	B724797	B724812	B724819	B724820	B724825	B724832	B724844	B724847	B724852	B724853	B724855	B724860
Shipment No.	1.	IP	11P	IP	1.	IP	1.	11	11	1.	IP	16	11	11	11	1.	1P	11	1.	11	dI

		Result	s of Prescr	Results of Prescreen Testing			Results	Results of Primary Screen Testing ^b	y Screen T	esting	
Shipment No.	Prescreen No.	YF	VE	Ħ	AVS.	VR	æ	VE	Ħ	SF	M
1P	B724863	65	1	2	6275	1	0	0	0	0	N.
1P	B724866	16	1	58	6704	0	0	0	0	0	0
IP	B724871	0	-	51	6276	0	0	0	0	0	MR
1P	B724885	24	1	65	9029	0	0	0	0	0	0
IP	B724886	25	-	19	6277	0	0	0	0	0	NR
4P	B724983	28	-	8	7411	0	0	0	0	0	0
4P	B724984	21	1	0	7412	0	0	0	0	0	0
4P	B724990	12	1	9	7413	0	0	0	0	0	1,2
IP	B724898	88	1	39	9029	1	0	0	0	0	0
7P	B848622	3	1	29	8254	1	0	0	0	0	0
7P	B848628	0	2	23	8255	1	0	0	0	1	.0
111	B848631	1,5	1,16	13,45	9270	0	0	1	2	0	0
11P	B848633	0	5	22	9271	0	0	0	2	0	0
111	B848634	7	1	13	9272	0	0	0	0	0	0
11P	B848635	0	0	17	9273	0	0	0	0	0	0
11P	B848649	6	40	19	9274	1	0	0	0	1	2
11P	B848650	4	8	13	9275	0	0	0	0	0	0
111	B848653	0,41	62,6	61,76	9276	0	1	7	. 2	2	0
11P	B848654	6,0	2,9	22,21	7776	0	0	0	0	0	0
111P	B848656	0,5	6,1	11,28	9278	0	0	1	0	0	0
111P	B848657	0	0	15	9279	0	0	0	0	0	0

Results of Prescreen Testing

Results of Primary Screen Testing^b

0					
0					
	0 0 0 0 0				
0					
9281	9282 9283 9284 9285	9282 9283 9285 9286 9287 9288	9282 9283 9284 9285 9286 9289 9290	9282 9283 9284 9285 9286 9286 9290 9290 9293	9282 9283 9284 9286 9286 9290 9290 9293 9294 9295
111	18 82,36 19,19				
2 - 0	2,2	3,10 3,10 1,2 1,2	2,2 3,10 1,2 1 1 2	2,2 3,10 1,2 1 1 1 2 0,0 0,0	2,2 3,10 1,2 1 1,2 1 0,0 0,0 2 2 2
0 0 0	8,0	0,3	8,0 8,0 0 0 0 0 0	0,3 0,8 0 0 0 0 0,18 5	0,3 0,8 0 0 0 0,18 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0
B848660 B848662 B848663	B848669 B848670	B848669 B848670 B848671 B848673 B848676	B848669 B848671 B848673 B848676 B848678 B848679 B848681	B848669 B848671 B848671 B848673 B848676 B848679 B848691 B848691	B848669 B848671 B848673 B848678 B848678 B848679 B848699 B848700 B848700
88 88 B8	B8	88 88 88 88 88 88 88 88 88 88 88 88 88	8	88 88 88 88 88 88 88 88 88 88 88 88 88	B B

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		Result	s of Prescr	Results of Prescreen Testing			Results	of Primar	Results of Primary Screen Testing ^b	esting	
Shipment No.	Prescreen No.	YF	VE	.	AVS No.	YF	JE	VE	М	SF	W
111	B848720	1	4	24	9301	1	0	1	1	0	0
111	B848722	3,2	1,1	3,11	9302	1	0	0	1	0	0
111	B848724	2	0	31	9303	2	1	0	1	0	0
111P	B848725	5,5	1,1	3,23	9304	0	1	0	1	0	0
11P	B848727	2,19	3,73	5,18	9305	1	0	1	1	0	0
11P	B848728	0,0	1,1	3,18	9306	1	0	0	1	0	0
11P	B848729	0	1	35	9307	1	0	0	1	0	0
111	B848730	0,9	4,6	1,4	9308	0	0	1	1	0	0
111	B848731	4,0	9,5	4,6	9309	0	0	0	1	0	0
11P	B848732	0	\$	34	9310	1	0	1	1	0	0
IIP	B848733	0,0	0,1	8,12	9311	0	0	0	1	0	0
IIP	B848734	0,0	0,1	3,23	9312	0	0	0	0	0	0
11P	B848735	3,1	0,0	11,16	9313	0	0	0	1	0	0
111P	B848736	2,1	6,18	9,19	9314	0	0	1	1	0	0
11P	B848737	6,5	0,1	1,8	9315	1	0	0	1	0	0
11P	B848738	1,0	1,3	9,26	9316	1	0	0	1	0	0
IIP	B848739	1	14	35	9317	2	0	1	2	0	0
111	B848740	5,0	2,9	7,26	9318	1	0	1	1	0	0
111	B848741	8	9	36	9319	1	0	0	2	1	0
111	B848742	3,1	6,0	0,5	9320	1	0	0	0	1	0
IIP	B848745	0	2	25	9321	0	. 0	1	0	0	0

Results of Prescreen Testing

W	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0,0	0	0
SF	0	2	0	0	0	2	2	0	0	0	1	0	0	0	0	1	2	0	0	1	1
FT	2	1	2	2	0	2	0	0	0	0	0	0	0	0	0	1	0	2	0	0	2
VE	1	0	0	0	0	1	0	0	0	0	0	0	0	0	0	0	0	1	0	0	1
3f	1	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
YF	1	0	1	0	0	1	0	1	0	0	0	0	0	0	0	0	. 0	0	0	0	0
AVS No.	9322	9323	9324	9325	9326	9327	9328	9329	9330	9331	9332	9333	9334	9335	9336	9337	9338	9339	9340	9341	9342
R	43,46	35	42	88	23	74,86	22,54	20	8,17	28	6,0	0,4	17	18	77	31	5,41	39	0,43	0,7	21,16
VE	4,9	4	1	5	4	2,22	81,6	3	0,0	1	2,0	1,1	1	3	1	1	4,21	7	9,0	0,27	2,40
YF	0,0	0	0	0	0	5,50	0,0	3	0,0	*	7,5	0,0	1	1	3	0	0,0	16	41,0	22,0	0,11
Prescreen No.	B848747	B848748	B848749	B848750	B848751	B848752	B848753	B848755	B848757	B848767	B848771	B848774	B848782	B848788	B848790	B848792	B848793	B848794	B848795	B848796	B848797
Shipment No.	111P	11P	111P	111	111	111	111	IIP	IIP	111	111	111	111P	111P	11P	11P	111	111	111	. 11P	IIP

Results of Prescreen Testing*

W	•	0	0	0	0	0	•	0	0	0	0	0	0	0		0	0				
Λ																		_			_
SF	0	0	0	0	0	0	0	0	0	0	0	2	0	0	0	0	0	•	•	1	0
M	0	0	0	0	. 0	0	0	1	0	0	0	1	0	0	0	0	0	0	0	0	0
VE	0	1	0	0	0	1	1	2	1	1	1	0	0	1	1	1	1	2	0	1	-
JE	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
YF	0	0	0	0	0	0	0	1		1	0	0	0	1	1	1	1	0	1	1	0
AVS No.	9343	9344	9345	9346	9347	9348	9349	9350	1326	9352	9353	9354	9355	9356	9357	9358	9359	9360	9361	9362	8986
PT.	0,17	2	6	99'0	4,16	1,2	0,17	1,19	3,55	2,40	25	7	32	3,43	8,55	0,27	19,21	0,26	25	23	2
VE	9,1	3	21	0,12	6,1	1,14	3,44	2,12	1,42	0,32	1	- 11	10	4,21	8,25	2,12	2,24	1,10	2	25	2
YF	0,4	24	5	0,0	0,0	0,0	6,17	0,0	0,1	0,20	0	2	0	6,0	0,0	2,0	0,0	1,0	0	3	0
Prescreen No.	B848798	B848800	B848801	B848804	B848805	B848808	B848809	B848810	B848811	B848814	B848816	B848820	B848838	B848839	B848841	B848843	B848845	B848848	B848861	B848864	R848866
Shipment No.	IIP	111	IIP	IIP	IIP	9P	IIP	IIP	IIP	111P	111P	IIP	IIP	12P	12P						

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3		0	0		•								0	0	0	0	0	2	0	0	•
>	Ľ	_			_								_	_	_	_	_				_
SF	2	1	0	0	0	0	0	1	0	0	0	0	0	0	0	0	0	0	0	0	•
PT	0	0	0	0	0	0	0	0	0	0	0	0	1	1	1	1	1	1	1	1	-
JE VE PT SF	0	1	2	0	0	0	1	•	1	0	•	1		0	1	1	0	2	0	0	-
JE	0	0	0	0	0	0	0	•	0	0	0	0	0	0	0	0	0	0	0	0	•
YF	2	-	0	0	0	0	1	1	1	•	-	0	0	0	0	0	0	0	0	0	•
AVS No.	9364	9365	9366	9367	9368	9369	9370	9371	9372	9373	9374	9375	9376	9377	9378	9379	9380	9381	9382	9383	9384
VE PT	15	32	34	39	23	40	56	53	57	45	•	28	23	35	30	29	48	45	47	4	61
VE	20	32	2	0	1	6	1	0	0	2	12	15	91	6	12	,	9	7	9	=	80
YF	9	5	5	3	0	*	2	0	0	•	•	0	•	•	0	10	0	0	0		18
Prescreen No.	B848869	B848870	B848873	B848874	B848876	B848879	B848880	B84881	B848882	B84883	B848892	B848893	B848895	B848896	B848897	B848899	B848900	B848901	B848903	B848904	B848905
Shipment No.	12P	12P	12P	12P	12P	12P	12P	12P	12P	12P	12P	12P	12P	12P	12P						

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Shipment	Prescreen	YF	VE	Z	AVS	· YF	æ	VE	Z	SF	\$
No.	No.				No.		7				
12P	B848906	0	12	35	9385	0	0	0	1	1	0
12P	B848907	0	30	52	9386	0	0	0	1	0	•
12P	B848909	•	9	31	9387	0	0	1	1	0	0
12P	B848911	0	25	79	9388	1	0	1	2	•	1
12P	B848913	0	3	**	9389	0	0	7	2	0	-
12P	B848914	61	0	23	9390	0	0	0	0	0	0
12P	B848916	18	0	7	9391	1	0	0	2	1	0
12P	B848917	22	0	0	9392	0	0	0	0	1	0
12P	B848920	1	S	31	9393	0	0	1	2	1	1
12P	B848921	1	3	48	9394	0	0	8	2	3	0
12P	B848924	13	-	3	9395	1	0	0	0	0	0
12P	B848926	1	2	99	9396	1	0	0	2	0	0
TP TP	B848985	0	•	47	8256	0	0	0	1	0	0
TP	B848986	6	3	88	8257	1	0	0	1	0	0
7.	B848989	0	0	199	8258	0	0	1	1	0	0
7.	B848990	0	•	53	8259	1,1	0,0	0'0	2,2	0,0	0
7.	B848991	•	1	40	8260	1,1	0,0	1,1	2,2	0,0	•
TP TP	B848992	12	3	4	8261	2,2	0,0	1,1	1,2	1,0	•
Ф.	B849022	13	18	88	8233	0,1	0,0	0,0	2,2	0,0	0
6P	B849035	91	6	99	8234	1,1	0,0	1,1	2,2	0,0	۰
GP	B849037	=	9	15	8235	0	0	ı	•	•	•

Results of Prescreen Testing*

		Result	s of Prescr	Results of Prescreen Testing			Results	Results of Primary Screen Testing ^b	Screen T	esting
Shipment No.	Prescreen No.	YF	VE	Z	AVS No.	YF	3	VE	M	SF
4P	GRP19424	3	1	27	7418	0,0	0,0	0,0	3,0	3,2
4P	GRP19425	0	1	13	7419	0	0	0	0	1
4P	GRP19427	0	1	12	7420	0	0	0	0	0
4P	GRP19432	0	-	10	7421	0	0	0	1	0
4P	GRP19433	0	1	18	7422	0	0	0	. 0	0
4P	GRP19434	15	1	14	7423	0	0	0	0	1
4P	GRP19435	9	1	25	7424	0,2	0,0	0,0	2,2	1,0
4P	GRP19437	20	1	5	7425	0	0	0	0	0
4P	GRP19438	0	-	14	7426	0	0	0	0	0
4P	GRP19439	23	-	9	7427	1,0	0,0	0,0	0,1	2,1
4P	GRP19440	15	1	6	7428	1,1	0,0	0,0	0,0	0,1
4P	GRP19441	11	1	27	7429	0,0	0,0	0,0	0,1	2,0
4P	GRP19442	20	1	20	7430	1,0	0,0	0,0	0,2	3,1
4P	GRP19446	7	1	9	7431	0	0	0	0	1
4P	GRP19447	6	1	20	7432	0	0	0	0	1
4P	GRP19448	9	1	8	7433	2,2	2,2	0,0	0,0	0,0
4P	GRP19449	5	1	6	7434	0,2	0,0	0,0	1,2	2,2
4P	GRP19450	1	1	80	7435	0,1	0,0	0,0	0,1	1,1
16P	GRP19457	10	23	32	11131	2	1	2	7	2
16P	GRP19458	14	26	31	11132	1	0	0	1	0
16P	GRP19459	13	80	83	11133	2	0	0	3	2

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Results of Prescreen Testing*

W	NT	T.N	N.	TN	IN	NT	NT	NT	NT	NT	IN	IN	IN	F.	N.	N.	¥	N.	N.	N.	r.
SF	3	0	0	1	1	1	1	1	0	0	1	0	0	1	NT	0	1	0	0	0	0
M	0	2	0	0	1	2	2	0	0	0	2	0	0	0	NT	0	2	0	1	0	1
VE	1	0	1	0	1	1	1	0	0	1	2	0	0	0	NT	1	2	1	0	1	0
JE	0	0	0	0	0	0	0	0	0	0	0	0	0	0	NT	0	0	0	0	1	0
YF	0	1	0	1	0	1	0	0	0	1	1	0	0	0	NT	1	0	0	0	1	0
AVS No.	11134	11135	11136	11137	11138	11139	11140	111141	11142	11143	11144	11145	11146	11147	11148	11149	11150	11151	11152	11153	11154
Ę	72	52	17	16	5	48	23	41	46	23	35	2	39	46	46	35	52	45	50	28	31
VE	30	3	61	61	1	5	15	15	2	27	12	13	12	30	30	17	63	49	2	2	21
YF	0	11	13	0	27	7	7	0	0	42	1	21	2	4	4	0	0	0	0	0	1
Prescreen No.	GRP19462	GRP19463	GRP19464	GRP19465	GRP19467	GRP19469	GRP19470	GRP19471	GRP19472	GRP19473	GRP19474	GRP19479	GRP19480	GRP19481	GRP19481	GRP19486	GRP19490	GRP19491	GRP19492	GRP19493	GRP19501
Shipment No.	16P																				

Results of Prescreen Testing*

		NCOU	2011100	ाइ जा हा इस हिला			MOUIN	Accounts of Finnary Serven Learning	od cell I	Sime	
Shipment No.	Prescreen No.	YF	VE	F	AVS No.	YF	JE	VE	PT	SF	W
	GRP19502	27	41	28	11155	0	1	2	2	0	NT
	GRP19509	0	2	35	11156	0	0	1	1	0	NT
	GRP19510	7	8	33	11157	0	0	1	0	0	N.
	GRP19511	0	12	57	11158	0	0	. 1	0	0	NT
	GRP19512	2	4	29	11159	0	0	0	0	0	NT
	GRP19514	15	1	12	11160	0	0	1	0	0	NT
	GRP21185	2	-	43	7436	0	0	0	0	0	0
	GRP21186	-		11	7437	0	0	0	0	0	0
	GRP21188	11	43	34	111161	0	0	0	1	0	NT
	GRP21189	13	0	32	11162	0	0	0	1	0	N
	GRP21190	13	0	6	11163	0	0	0	0	0	NT
	GRP21191	14	0	21	11164	0	0	0	0	0	N
	GRP21194	11	. 4	45	11165	0	0	0	0	0	N
16P	GRP21195	5	3	50	11166	1	0	0	2	0	NT
	GRP21197	1	30	47	11167	0	0	0	0	0	N
	GRP21199	8	23	2	11168	0	0	2	1	1	M
	GRP21201	11	0	20	11169	0	0	0	0	0	NT
	GRP21205	3	70	25	11170	0	0	1	0	1	Į.
	GRP21207	30	24	47	11171	1	0	0	0	0	N
	GRP21214	0	36	16	11172	0	0	1	0	0	NT

- a = Maximum percent reduction in CPE observed in the assay.
- = Explanation for Abbreviations/Numbers (Each number represents one test):
- Active at the 95% level (% reduction in viral CPE of \geq 95% at one or more concentrations). II
- Active at the 50% level (% reduction in viral CPE of \geq 50% to \leq 94% at one or more concentrations). 11
- Active at the 25% level (% reduction in viral CPE at ≥25% to ≤49% at one or more concentrations). II
- 0 = Inactive (<25% reduction in viral CPE at all concentrations).
- NT = Not Tested.
- NR = Not Required.

Note:

Symbols are separated by commas where more than one test was performed (i.e. "1,0" designates two tests, one was active at the 25% level and the second test was inactive).

4.3. Antiviral Evaluations In Vivo:

4.3.1. Pichinde Virus in MHA Hamsters:

A total of 19 compounds were assessed for efficacy against Pichinde in vivo. These compounds are listed in Table 35. Of the compounds evaluated, only 2 (AVS 01 and AVS 206) had significant activity against Pichinde virus in vivo. AVS 01 is ribavirin, which has broad spectrum activity against RNA viruses. The compound was able to decrease the mortality and increase the average day of death in virus-infected hamsters. The antiviral efficacy of ribavirin was dose-dependent with the greatest efficacy demonstrated at a dose of 150 mg/kg/day. Dose levels above this were toxic to the hamsters.

AVS 206, an analog of AVS 01, was also active against Pichinde virus in vivo. At 100 mg/kg/day AVS 206 increased the average day of death; however, there was no significant reduction in the mortality. The results of this assay are presented in Table 36. A second assay of this compound was conducted at a dose level of 200 mg/kg/day administered on a qld x 12 days schedule with the first dose administered on the day preceding virus challenge. In this study, AVS 206 reduced mortality and prolonged the average day of death significantly (Table 37). These results were encouraging and led to an additional study with AVS 206. For this, AVS 206 was administered at doses of 1000, 500, 250 and 200 mg/kg/day on a qld x 10 days schedule beginning on the day before virus challenge. AVS 01 (ribavirin) was administered to hamsters at 500, 200 and 100 mg/kg/day on the same treatment schedule for comparison. As shown in Table 38, AVS 206 was toxic at 1,000 and 500 mg/kg/day while the 250 and 200 mg/kg/day dose levels reduced mortality markedly. In contrast, AVS 01 was toxic at doses of 200 mg/kg/day and higher. These results indicate that the ribavirin analog, AVS 206, may be less toxic than ribavirin although no enhancement in antiviral activity was noted with this carboxamidine analog of ribavirin.

AVS 1046 was considered a candidate for further testing against arenavirus infections. As shown in Table 39, treatment with AVS 1046 prolonged the ADD in Pichinde-infected hamsters in a dose-dependent manner. Since the compound was well tolerated at the highest test dose (100 mg/kg/day) further testing at higher doses was recommended; however, there was no drug available for the studies.

Of the 19 compounds evaluated against Pichinde virus only 2 demonstrated good potential for treatment of arenaviral infections. These compounds, AVS 001 and AVS 206, significantly reduced virus induced mortality. Other compounds (AVS 79, 94, 111, 167, 332, 346, 925, 1046 and 1250) which were tested at concentrations below the maximum tolerated dose were not available in quantities sufficient for further evaluations. These compounds should be re-tested at higher dose levels when sufficient compound quantities become available.

Several problems are present in the Pichinde virus infection model. First, the virus only produced disease in inbred strain MHA hamsters. These hamsters are only available from 1 commercial source. The disease is more severe in the female than in the male hamster which must be considered during design of the experimental protocol. Thirdly, the viral infection is complicated by gastrointestinal disease which is presumed to be of bacterial origin. Attempts to treat the gastrointestinal disease with oral tetracyclines did not prove effective in our hands. Studies with AVS 01 indicate that the greatest antiviral efficacy was achieved if treatments were administered on a 10 day schedule rather than a 7 day schedule. The critical timepoint for administration of AVS 001 appears to be between days 2 and 6 post-infection. If this model is pursued for further antiviral drug testing, we recommend that tissue virus titers be evaluated in addition to monitoring mortality and ADD.

Table 35

Compounds Assessed Against Pichinde Virus In Vivo

AVS#	Activity	Maximum tolerated dose	Recommendation
01	yes	150 mg/kg/day	positive control drug
52	no	10 mg/kg/day	inactive
79	no	>100 mg/kg/day	retest at higher doses
94	no	>100 mg/kg/day	retest at higher doses
111	no	> 150 mg/kg/day	retest at higher doses
148	no	10 mg/kg/day	inactive
167	no	>32 mg/kg/day	retest at higher doses
206	yes	250 mg/kg/day	pursue further
215	no	32 mg/kg/day	inactive
253	no	32 mg/kg/day	inactive
272	no	32 mg/kg/day	inactive
332	no	> 100 mg/kg/day	retest at higher doses
346	no	>60 mg/kg/day	retest at higher doses
347	no	<5 mg/kg/day	retest at higher doses
360	no	10 mg/kg/day	inactive
646	no	10 mg/kg/day	inactive
925	no	> 100 mg/kg/day	retest at higher doses
1046	possible	> 100 mg/kg/day	retest at higher doses
1250	no	>100 mg/kg/day	retest at higher doses

Table 36
ANTIVIRAL EPFICACY OF AVS-206

	7	
	×	
	q1d	
Pichinde	Schedule:	Day 0
Virus:	Dosage Sch	Startir

Starting: Day O	UNINFECTED	INFECTED	GMTD	N P
Untreated	0/5	$10/10 (10.1 \pm 1.1)^{b}$	10.1	6.0
PBS	9/2	10/10 (10.8 ± 3.3)	10.4	1.0
100 mg/kg AVS 206	0/5	8/10 (15.3 ± 2.4)	17.1	1.6
10 mg/kg AVS 206	0/2	9/10 (13.1 ± 2.5)	13.9	* e. c
3.2 mg/kg AVS 206 100 mg/kg Ribavirin	0/5	7/10 (16.3 ± 1.8)	10.3	1.8
a * dead/* treated				

dead/* treated ADD = $\sum [(day of death) \times (* dead that day)] + S.D.$ total number of dead

GMTD = geometric mean time to death

VR = GMTD of Experimental GMTD of PBS Control

Table 37

RESULTS OF RETESTING AVS-206

Dosage Schedule: q1d x 12 Starting: Day -1 Virus: Pichinde

	UNINFECTED	INPECTED	GMTD	K _d
Untreated	0/58	10/10 (9.5 ± 1.3) ^b	9.4	1.0
PBS	0/5	$10/10 (9.7 \pm 0.9)$	9.7	1.0
200 mg/kg AVS 206	9/2	$4/10^{6} (18.8 \pm 1.5)^{f}$	23.8	2.5
100 mg/kg Ribavirin	$1/5 (18.0 \pm 0.0)$	$3/10^{gh}(21.7 \pm 4.5)^{1}$	25.8	2.7
Virus Titration				
1 X 107 pfu/hamster				
1 X 106 pfu/hamster		5/5 (8.6 + 1.1)		
1 X 105 pfu/hamster				
1 X 104 pfu/hamster				
1 X 103 pfu/hamster		5/5 (9.4 + 1.1)		
1 X 102 pfu/hamster		5/5 (10.2 + 1.9)		
1 X 101 pfu/hamster		5/5 (9.8 ± 1.3)		
a # dead/# treated	# dead/# treated			
ADD = 2 ((day of	death) x (# dead that day	(1) + S.D.		

total number of dead

Pishers' Exact Test comparing % mortality ribavirin & PBS p = .0015 Fishers' Exact Test comparing % mortality ribavirin & AVS 206 p = 0.5Pishers' Exact Test comparing % mortality AVS 206 & PBS VR = GMTD of Experimental GMTD of PBS Control p = .000003T-test of ADD p = <10-6T-test of ADD

GMTD = geometric mean time to death

Table 38

Antiviral Efficacy of AVS-206 and Ribavirin (AVS-01)

Virus: Pichinde

Treatment: q1d x 10 days starting on the day preceding virus challenge

		No. Dea				No. In				GMTD*	<u>VR</u> ^e
Untreated		0/10				10/10	(9.8	±	0.9)	9.8	NA
PBS		0/10				10/10	(9.0	±	1.3)	8.9	1.0
1000 mg/kg			(5.0			10/10	(5.0			4.9	0.6
500 mg/kg	AVS-206	4/4	(8.5	±	0.6)	10/10	•		1.5)	8.3	0.9
250 mg/kg	AVS-206	0/5				1/104	(23.0)	±	0.0)*	27.5	3.1
200 mg/kg	AVS-206	0/5				2/10 ^r	(25.0	±	2.8)*	27.4	3.1
1000 mg/kg	AVS-01	5/5	(3.6	±	0.5)		No	t de	one		
500 mg/kg.	AVS-01	5/5	(5.2	±	0.4)	10/10	(5.7	±	1.3)	5.6	0.6
250 mg/kg	AVS-01	5/5	(7.6	±	2.4)		No	t de	one		
200 mg/kg	AVS-01	3/5	(10.7	±	1.5)	8/10	(12.8	±	4.4)b	14.4	1.6
100 mg/kg	AVS-01	0/5				3/10	(18.7	±	8.1) ^j	24.4	2.7
Virus challe	nge										
1 x 10° pfu		NA				5/5	(8.4	±	0.9)	8.4	NA
1 x 10 ⁵ pfu		NA				5/5	(9.0	±	1.2)	8.9	NA
1 x 10 ⁴ pfu		NA				5/5	(8.2	±	0.4)	8.2	NA
1 x 10 ³ pfu		NA				5/5			0.5)	9.4	NA
1×10^2 pfu		NA				5/5			0.7)	9.0	NA
10 pfu		NA				5/5			1.6)	9.9	NA

a. No. in parentheses = Average Day of Death \pm 1 Standard Deviation (ADD \pm SD)

ADD = $\sum [(\text{day of death}) \times (\text{number dead that day})]$ total number of dead

b. GMTD = Geometric Mean Time to Death

GMTD = $x\sqrt{n_1 \cdot n_2 \cdot n_x}$ where n = day of death x = total number of animals All survivors are calculated as dying on Day 28

- c. VR = GMTD of Experimental/GMTD of Diluent-Treated Control
- d. Fisher's Exact Test comparing mortality of PBS-treated vs drug treated $p = 5.9 \times 10^{-5}$
- e. Student's T-test comparing ADD of PBS-treated vs drug treated $p = 3.5 \times 10^4$
- f. Fisher's Exact Test comparing mortality of PBS-treated vs drug treated $p = 3.6 \times 10^{-4}$
- g. Student's T-test comparing ADD of PBS-treated vs drug treated p < 104
- h. Student's T-test comparing ADD of PBS-treated vs drug treated p = .02
- i. Fisher's Exact Test comparing mortality of PBS-treated vs drug treated $p = 1.6 \times 10^{-3}$
- j. Student's T-test comparing ADD of PBS-treated vs drug treated p = .002

ANTIVIRAL EFFICACY OF AVS-1046 Table 39

Virus: Pichinde Dosage Schedule: q1d x 7 Starting: Day 0

	UNINFECTED	INFECTED	GWTD ^C	K.
Untreated	0/5 ^a	10/10 (10.5 ± 1.8) ^b	10.4	1.0
0.4% CMC in PBS	9/0	10/10 (10.8 ± 3.5)	10.4	1.0
100 mg/kg AVS 1046	0/5	10/10 (13.7 + 2.9)	13.4	1.3
32 mg/kg AVS 1046	0/5	9/10 (12.3 + 2.7)	13.1	1.3
10 mg/kg AVS 1046	0/5	10/10 (12.4 + 4.7)	11.8	1.1
3.2 mg/kg AVS 1046	0/2	9/10 (10.9 ± 2.3)	11.8	1.1
100 mg/kg Ribavirin	0/5	$7/9 (15.9 \pm 5.3)$	17.7	1.7
b ADD = \(\sum_{\begin{subarray}{cccccccccccccccccccccccccccccccccccc	d of death) x (* dead that day)] + S.D.	(V)] ± S.D.		
total numbe	number of dead			

GMTD = geometric mean time to death ပ

VR = GMTD of Experimental GMTD of PBS Control

4.3.2. Venezuelan Equine Encephalomyelitis Virus (VE):

A total of 6 compounds were received for testing against VE. These compounds are listed in Table 40. None of the compounds had significant activity against VE induced mortality; however, 5 of the 6 compounds were not available for testing at maximum tolerated dose levels. Thus, it is possible that these compounds may show activity against VE if higher dose levels are tested.

Because of the problem of limited compound availability, we initiated studies to develop a target organ model for VE. For these studies, a series of experiments were conducted to determine the lethal doses of VE following intracranial challenge. The cumulative results of these experiments are presented in Table 41. From this a challenge inoculum of 0.03 ml of a 10^{-8.3} dilution of the VEE virus stock was selected for a chemotherapy study. We had identified a compound, 3 nitro-3-deazauridine (3N-3DU), from other studies which had potential as a chemotherapeutic agent against alphaviruses.

The effect of a single intracranial dose of 40 mg of 3N-3DU/kg on VEE infection was assessed. Mice (10/group) were challenged intracranially with 0.03 ml of a 10^{-8.3} dilution of VEE stock virus and treated 6 or 8 or 10 hours later with 0.03 ml of PBS or 3N-3DU intracranially. Mice receiving 40 mg/kg 3N-3DU at 8 hr post virus challenge had a mortality rate of 25% compared to 70% in the PBS treated group. In addition, the ADD for the 3N-3DU treated group was significantly increased (p = 0.049). These results are shown in Table 42. Thus, the target organ approach may be a viable alternative for assessing compounds available in limited quantities. Unfortunately, changes in the contract workscope prevented us from pursuing development of this potential model system.

Several recommendations can be made from our studies with this VEE animal model system. First, a less severe challenge virus would be preferable to the Trinidad donkey strain used in our studies. This virus, even at low inoculum doses, is a severe challenge in mice. Since most compounds are available in very limited quantities and little, if any, is known about their pharmacokinetics it is difficult to select an appropriate treatment dose and schedule. Therefore, it is difficult to demonstrate antiviral efficacy in preliminary studies. A second possible improvement would be to quantitate virus titers in the tissues rather than simply monitoring mortality and ADD. This may provide better evidence for antiviral efficacy. Thirdly, the target organ model may be a viable preliminary screening tool for assessing potential chemotherapeutic efficacy. Positive results from this system could be used to identify lead compounds for re-synthesis.

Table 40

Compounds Received for Testing Against Venezuelan Equine Encephalomyelitis Virus

AVS#	Activity	Maximum tolerated dose	Recommendation
79	no	100 mg/kg	inactive
206	no	>1,000 mg/kg	retest at higher doses
272	no	>100 mg/kg	retest at higher doses
347	no	>32 mg/kg	retest at higher doses
360	no	>100 mg/kg	retest at higher doses
646	no	> 100 mg/kg	retest at higher doses

Table 41

Cumulative Results of Intracranial Challenge with Venezuelan Equine
Encephalomyelitis Virus in Mice

Challenge Dose (0.03 ml)	#dead challenged	% Mortality
PBS	0/29	0%
10 ⁻⁶ dilution of VEE	10/10	100%
10 ⁻⁷ dilution of VEE	10/10	100%
10 ^{-7.1} dilution of VEE	20/20	100%
10 ^{-7.4} dilution of VEE	20/20	100%
10 ^{-7.7} dilution of VEE	18/20	90%
10 ⁻⁸ dilution of VEE	19/30	63 %
10 ^{-8.3} dilution of VEE	12/20	60%
10 ^{-8.6} dilution of VEE	6/19	32%
10 ^{-8.9} dilution of VEE	0/10	0%
10 ⁻⁹ dilution of VEE	2/10	20%

Table 42
Mortality in VEE Challenged Mice Receiving a Single Dose of 3N-3DU

Treatment Groups	No. Dead/No. Treated	ADD + 1 SD
Uninfected Mice		
Untreated control	0/5	N/A
Sham-infected IC	0/5	N/A
Sham-infected + PBS IC	0/5	N/A
Sham-infected + 40 mg/kg 3N-3DU	IC 0/4	N/A
Virus-Infected Mice		
Untreated control	2/5	6.5 +/- 0.7
Placebo (PBS)	4114	
6 hrs	6/10	6.3 + -0.8
Placebo (PBS)		
8 hrs	7/10	5.4 +/- 0.5
Placebo (PBS)		
10 hrs	5/10	5.8 +/- 1.5
40 mg/kg 3N-3DU		
6 hrs	6/10 (.68)	5.8 +/- 0.8
40 mg/kg 3N-3DU		
8 hrs	2/8 (.077)	6.5 +/- 0.7 (.049)
40 mg/kg 3N-3DU		
10 hrs	5/8 (.56)	6.5 + -1.2 (.52)

4.3.3. Japanese Encephalitis Virus (JE):

During the course of this contract, extensive attempts were made to establish a reproducible JE virus model in outbred Swiss mice with the Nakayama strain of JE. The only reproducible challenge identified with this virus strain in outbred Swiss mice was following intracerebral inoculation. Since a less severe challenge was desirable, we began studies with the Beijing strain of JE in inbred C57Bl/6 mice. These studies resulted in development of a lethal JE virus model. The challenge inoculum, prepared by serial in vivo passage through the CNS tissues of 4 week-old C57Bl/6 mice, was reproducibly lethal following intraperitoneal inoculation into C57Bl/6 mice.

A total of 13 compounds was received for testing against JE in vivo. These compounds are listed in Table 43. Of these, 7 had demonstrable activity against JE in vivo. These compounds were AVS #360, 361, 2811, 2812, 2979, 2980 and 5587.

As can be seen in Table 44, AVS 360 demonstrated some antiviral efficacy as the mortality rate in treated mice was 70% compared to 100% in the diluent-treated controls. The ADD of AVS 360 treated mice was not prolonged compared to the diluent-treated controls. Further studies with alternate dose, route and treatment schedules may provide improved antiviral efficacy.

A second assay of AVS 360 was conducted at dose levels of 300 and 150 mg/kg/day on a qld x 7 days schedule. The results, shown in Table 45, indicate a possible antiviral effect in that the mortality was slightly decreased. Since the maximum tolerated dose was not achieved it is possible that greater activity might be demonstrable. Additional testing of this compound at higher doses on alternate treatment schedules is recommended.

AVS 361 was initially tested at doses of 6, 4 and 2 mg/kg/day administered qld x 7 days starting the day preceding virus challenge. As shown in Table 46, AVS 361 was toxic at 6 mg/kg. At 4 mg/kg, the compound completely suppressed virus-induced mortality while the 2 mg/kg dose level did not significantly suppress mortality. However, the 2 mg/kg dose level did prolong the ADD significantly. A second assay with this compound confirmed its antiviral activity. The results, shown in Table 47, vary slightly from the first experiment with regard to the toxic dose level. In the second study, AVS 361 was tolerated at 6 mg/kg and this dose significantly reduced mortality. At 4 mg/kg the ADD was significantly prolonged although the mortality was not significantly decreased. The variance in toxic dose levels between the two assays could have resulted from differences in the compound since separate shipments were used for the two assays. Another potential source of variance is the preparation of the dosing solutions. The conclusion from these studies is that AVS 361 has potential for anti-JE activity in vivo. Although the window between toxic and therapeutic doses is narrow, it may be possible to enhance the antiviral effect by altering compound formulations, administration routes and by synthesizing analogs of this compound.

AVS 2563 may have an effect against JE. As shown in Table 45, the highest dose level tested, 45 mg/kg/day, reduced mortality slightly. Since the maximum tolerated dose was not achieved it would be advisable to retest this compound at higher dose levels before it is excluded as inactive.

AVS 2811 was initially tested at 25, 12.5 and 6.25 mg/kg/day on a qld x 7 days schedule beginning on the day preceding virus challenge. As shown in Table 46, the 25 mg/kg/day dose produced 80% mortality in the toxicity control mice. However, the virus-infected mice receiving this dose had only 40% mortality compared to 100% mortality in the virus-infected diluent treated mice. At the 12.5 and 6.25 mg/kg dose levels the mortality was 80% in the virus-infected mice with a slight prolongation of the ADD. This compound was assessed in a second study at doses of 20, 10 and 5 mg/kg/day. In this study the drug diluent was HPC obtained from the NCI in contrast to 2% alcohol in saline used for the first study. As shown in Table 47, the 20 mg/kg dose level was uniformly toxic. However, the 10

mg/kg/day dose level significantly reduced virus induced mortality. The 5 mg/kg dose level was ineffective in reducing mortality. The studies differ slightly with regard to toxicity and effective doses. This may be the result of differences in (1) the compound (received in separate shipments); (2) the drug preparation or (3) the drug diluent used in the 2 assays. In conclusion, AVS 2811 has some activity against JE virus in vivo and further studies should be pursued.

AVS 2812 was assayed for activity against JE virus at doses of 6, 3 and 1 mg/kg/day on a qld x 7 days schedule beginning on the day preceding virus challenge. As shown in Table 46, the 6 mg/kg dose level resulted in 80% mortality in the toxicity control mice with a 56% mortality in the virus-infected mice. At 3 mg/kg, AVS 2812 was not toxic to the toxicity control mice and it reduced mortality significantly in the virus-infected mice. Similarly, the 1 mg/kg dose level also decreased mortality. A retest of AVS 2812 at the same dose levels confirmed the activity as shown in Table 47. Although the window between toxicity and efficacy is narrow, this compound and analogs of it should be pursued in further studies.

AVS 2979 was initially evaluated at doses of 200, 100 and 50 mg/kg/day which were toxic. The compound was tested at 30 and 15 mg/kg/day on a qld x 7 days schedule beginning the day preceding virus challenge. The 30 mg/kg dose administered subcutaneously reduced mortality to 40% compared to 80% in the diluent-treated control mice. The mortality was reduced to 30% when treatment with 15 mg/kg was administered subcutaneously. This same dose level administered intraperitoneally only reduced mortality to 60% compared to the diluent-treated control of 78%. In this study, the subcutaneous route was superior to the intraperitoneal route. These results, shown in Table 48, indicate potential activity for AVS 2979. Testing at doses of 10 mg/kg and lower did not provide evidence of antiviral efficacy. Further studies with this compound are recommended.

AVS 2980 was tested at doses of 40, 20 and 10 mg/kg/day on a qld x 7 days schedule beginning on the day preceding virus challenge. At 40 mg/kg, AVS 2980 was not lethally toxic; however, it did produce weight loss in the toxicity control mice. This dose level significantly reduced mortality in the virus-infected mice. At 20 mg/kg, AVS 2980 reduced mortality to 40% compared to 100% mortality in the diluent-treated virus-infected mice. No activity was present at the 5 mg/kg dose level. These results, shown in Table 47, indicate significant potential for AVS 2980 although the effective and toxic dose levels are not significantly different. Further studies with this compound and its analogs are recommended.

AVS 5587 was significantly effective in reducing JE virus-induced mortality. The toxicity control mice receiving 200 mg/kg/day did not suffer any detectable drug toxicity as they gained weight throughout the experimental observation period. The virus-infected mice receiving 200 mg/kg/day had a mortality rate of 10% which is significantly reduced from the 100% mortality occurring in the diluent-treated, virus-infected mice. The virus-infected mice receiving 100 mg/kg/day had a mortality rate of 20% which also represents a significant reduction. At 50 mg/kg, AVS 5587 reduced mortality to 10%. In addition to reducing the mortality rates these dose levels also significantly prolonged the ADD. Doses of 10 mg/kg and lower were not effective in altering virus-induced mortality. These results, shown in Table 49, indicate significant potential for AVS 5587 as a chemotherapeutic agent. The results of this assay should be confirmed and further dose, route, schedule studies should be conducted. Of all the compounds demonstrating anti-JE activity, this compound may be most desirable as the window between toxicity and efficacy is broader than that seen with the other compounds. Further studies of this compound and its analogs are strongly recommended.

Table 43

Compounds Received for Testing Against Japanese Encephalitis Virus In Vivo

AVS#	Activity	Maximum tolerated dose	Recommendation
111	no	>320 mg/kg/day	retest at higher dose levels
215	no	64 mg/kg/day	inactive
257	no	>320 mg/kg/day	retest at higher dose levels
272	no	>53 mg/kg/day	retest at higher dose levels
360	yes	>300 mg/kg/day	retest at higher dose levels
361	yes	6 mg/kg/day	evaluated for potential use
2563	possible	>45 mg/kg/day	retest at higher dose levels
2811	yes	12.5 mg/kg/day	evaluated for potential use
2812	yes	3 mg/kg/day	evaluated for potential use
2979	yes	30 mg/kg/day	evaluated for potential use
2980	yes	>40 mg/kg/day	evaluated for potential use
4113	no	>32 mg/kg/day	retest at higher dose levels
5587	yes	>200 mg/kg/day	evaluate for potential use

Table 44

Antiviral Evaluation of AVS-360 (JE Virus)

Treatment	No. dead/ No. uninfected	No. dead/ No. infected ^a	GMTD ^b	VRc
Compounds administered	subcutaneously qld x	7 days starting day -1		
Untreated	0/5	$10/10 \ (13.6 \pm 1.5)$	13.5	NA^d
0.4% CMC in PBS	0/5	$10/10 \ (12.3 \pm 2.8)$	12.1	NA
250 mg/kg AVS-360	0.5	$7/10 \ (12.0 \pm 2.5)$	15.2	1.3

^{*} ADD ± 1SD = Average Day of Death ± 1 Standard Deviation

ADD = \sum [(day of death) x (number dead that day)] total number of dead

GMTD =
$$x \sqrt{n_1 \cdot n_2 \cdot n_x}$$
 where $n = \text{day of death}$
 $x = \text{total number of animals}$
All survivors are calculated as dying on Day 28

VR = GMTD of Experimental/GMTD of Diluent-Treated Control

^b GMTD = Geometric Mean Time to Death

c VR = Virus Rating

^d NA = Not Applicable

Table 45 In Vivo Antiviral Efficacy of AVS-360 and AVS-2563 Against JE Virus

	No. Dead/		No. Dead/ No.		р	р
Treatment	Uninfected	ADD + SD	Infected	ADD + SD	value ^a	value ^b
Compounds admini	stered subcut	aneously qld x 7	days starting	<u>day -1</u>		
Untreated	0/5		10/10	12.1 <u>+</u> 4.9	0.5	0.21
0.4% CMC in PBS	0/5	1.2 <u>+</u> 0.5	9/10	14.3 <u>+</u> 1.5	-	-
AVS-360						
300 mg/kg	0/5		7/10	13.6 <u>+</u> 1.3	0.29	0.245
150 mg/kg	0/5		8/10	13.5 ± 1.3	0.5	0.244
Compounds admini	stered intrap	eritoneally q1d x	7 days starting	<u>day -1</u>		
0.4% CMC in PBS	0/5		10/10	13.4 <u>+</u> 2.6	-	-
AVS-2563						
45 mg/kg	0/5		7/10	13.9 <u>+</u> 2.2	0.11	0.706
30 mg/kg	0/5		10/10	13.1 + 1.7	1.0	0.759
15 mg/kg	0/5		9/10	13.0 ± 2.7	0.5	0.743

p = value using Fisher's Exact Test comparing mortality to diluent-treated control.
 p = value using Student's T-test comparing experimental ADD to diluent-treated control.

Table 46

Antiviral Efficacy of AVS-361, AVS-2811 and AVS-2812 Against JE Virus

		o. Dead/ Uninfected ^a	<u>GMTD</u> ^b		o. Dead/ . Infected ^a	<u>GMTD</u> ^b	VRc
Untreated	0/5			10/10	(14.0 ± 1.8)	13.9	NA
2% alc/saline	0/5			10/10	(12.8 ± 1.3)	12.7	1.0
AVS-361							
6 mg/kg	3/5	(2.7 ± 0.6)	6.8	7/10	$(4.0 \pm 2.4)^{d}$	6.5	0.5
4 mg/kg	0/5	(311 311)		0/10 ^e		28.0	2.2
2 mg/kg	0/5			8/10	$(16.3 \pm 1.9)^{f}$	18.0	1.4
AVS-2811							
25 mg/kg	4/5	(4.8 ± 3.1)	6.0	4/10 ^g	(10.5 ± 6.7)	16.1	1.3
12.5 mg/kg	0/5	/		8/10	(14.0 ± 1.8)	16.0	1.3
6.25 mg/kg	0/5			8/10	(13.5 ± 1.5)	15.6	1.2
AVS-2812							
6 mg/kg	4/5	(6.0 ± 1.4)	8.0	5/9h,i	$(6.4 \pm 3.1)^{i}$	15.8	1.2
3 mg/kg	0/5			2/10 ^k	$(17.5 \pm 7.8)^{1}$	25.2	2.0
1 mg/kg	0/5			6/10 ^m	$(15.7 \pm 3.4)^n$	19.5	1.5
Sham-infected	0/10						
4 x challenge	NA		NA	5/5	(13.4 <u>+</u> 1.1)	13.4	NA
2 x challenge	NA		NA	5/5	(12.2 ± 1.1)	12.2	NA
1 x challenge	NA		NA	5/5	(12.4 ± 2.7)	12.1	NA
0.5 x challenge	NA		NA	5/5	(13.4 ± 1.1)	13.4	NA

^a No. in parentheses = Average Day of Death + 1 Standard Deviation (ADD ± 1SD)

ADD = $\sum [(\text{day of death}) \times (\text{number dead that day})]$ total number of dead

GMTD =
$$x \sqrt{n_1 \cdot n_2 \cdot n_x}$$
 where $n = day \text{ of death}$ $x = total number \text{ of animals}$

All survivors are calculated as dying on Day 28

^b GMTD = Geometric Mean Time to Death

^c VR = GMTD of Experimental/GMTD of diluent-treated Control.

^d T-test of ADD vs. diluent-treated controls p < 10⁻⁶.

^e Fishers' Exact Test of exptal vs. control mortality $p = 5 \times 10^6$.

^f T-test of ADD vs. diluent-treated controls $p = 3.4 \times 10^4$.

⁸ Fishers' Exact Test of exptal vs. control mortality $p = 5.4 \times 10^3$.

h One animal sacrificed on day 28 due to head tilt not included in data.

i Fishers' Exact Test of exptal vs. control mortality p = 0.047.

^j T-test of ADD vs. diluent-treated controls $p = 7 \times 10^{-5}$.

k Fishers' Exact Test of exptal vs. control mortality p = 3.6 x 10⁻⁴.

T-test of ADD vs. diluent-trated controls p = .052.

m Fishers' Exact Test of exptal vs. control mortality p = 0.04.

ⁿ T-test of ADD vs. diluent-treated controls p = .031.

Table 47

Antiviral Efficacy of Selected AVS Compounds
Against JE Virus

Treatment	No. dead/ No. uninfected	No. dead/ No. infected ^a	<u>GMTD</u> ^b	VRc
Untreated	0/5	10/10 (12.2 ± 2.5)	11.9	NAd
2% alc/saline	0/5	10/10 (12.4 <u>+</u> 2.6)	12.1	NA
AVS-361				
6 mg/kg	0/5	$1/10^{e} (15.0 \pm 0.0)^{f}$	26.3	2.2
4 mg/kg	0/5	$7/10^g$ (15.4 \pm 2.0) ^h	18.4	1.5
2 mg/kg	$1/5 (28.0 \pm 0.0)$	$9/10 \ (11.6 \pm 1.4)$	12.5	1.0
AVS-2812				
6 mg/kg	$3/5 (7.0 \pm 5.0)$	10/10 (6.9 <u>+</u> 2.8) ⁱ	6.3	0.5
3 mg/kg	0/5	$1/10^{j} (15.0 \pm 0.0)^{k}$	26.3	2.2
1 mg/kg	0/5	$9/10 \ (15.1 \pm 1.2)^{1}$	16.0	1.3
NCI/HPC	0/5	· 10/10 (12.6 <u>+</u> 1.2)	12.6	NA
AVS-2811				
20 mg/kg	$5/5 (2.6 \pm 1.3)$	$10/10 (3.5 \pm 2.2)^{m}$	2.8	0.2
10 mg/kg	0/5	$4/10^{n} (13.3 \pm 1.0)^{o}$	20.7	1.6
5 mg/kg	0/5	9/10 (13.4 <u>+</u> 1.9)	14.4	1.1
AVS-2980				
40 mg/kg	0/5	$2/10^{p} (9.5 + 4.9)^{q}$	22.2	1.8
20 mg/kg	0/5	$4/10^{\rm r}$ (15.0 \pm 2.6)*	21.7	1.7
10 mg/kg	0/5	$9/10 (13.6 \pm 1.9)$	14.4	1.1

^a The numbers in parentheses are the ADD \pm SD = average day of death \pm 1 standard deviation

ADD = $\sum [(\text{day of death}) \times (\text{number dead that day})]$ total number of dead

b GMTD = Geometric Mean Time to Death

GMTD = $x \sqrt{n_1 \cdot n_2 \cdot n_x}$ where n = day of death

x = total number of animals

All survivors are calculated as dying on Day 28

c VR = Virus Rating

VR = GMTD of Experimental/GMTD of Diluent-Treated Control

^d NA = Not Applicable

^e Fisher's Exact Test $p = 5.95 \times 10^{-5}$.

f Student's t-test p = .36.

Fisher's Exact Test p = .105.

h Student's t-test p = .02.

i Student's t-test $p = 2.7 \times 10^{-4}$.

^j Fisher's Exact Test $p = 5.95 \times 10^{-5}$.

k Student's t-test p = .36.

Student's t-test p = .01.

^m Student's t-test = $< 10^{-6}$.

ⁿ Fisher's Exact Test $P = 5.4 \times 10^{-3}$.

O Student's t-test p = .35.

^p Fisher's Exact Test $p = 3.6 \times 10^{-4}$.

^q Student's t-test p = .06.

Fisher's Exact Test $p = 5.4 \times 10^{-3}$.

Student's t-test p = .03.

Table 48

In Vivo Testing of AVS-2979 Against JE Virus Challenge

Virus: JE

JE Experiment No. 38

Treatment: q1d x 7, starting Day -1

Treatment Group	No. Dead/ No. Uninfected	No. Dead/ No. Infected	ADD ± 1 SD ^a
Administered Subcutaneously	Ĺ		
Untreated	0/5	9/10	12.1 <u>+</u> 2.0
0.4% CMC	0/5	8/10	13.6 ± 1.8
30 mg/kg AVS-2979 15 mg/kg AVS-2979	0/5 0/5	4/10 b 3/10 d	14.3 ± 1.3 ° 16.0 ± 1.7 °
Administered Intraperitoneal	ly		
0.4% CMC	0/5	7/9	15.1 <u>+</u> 2.4
15 mg/kg AVS-2979	0/5	6/10 ^f	11.8 + 1.2 8

^a ADD + 1SD = Average Day of Death + 1 Standard Deviation

b Fisher's exact test versus diluent treated control

c Student's t-test versus diluent treated control

d Fisher's exact test versus diluent treated control

s Student's t-test versus diluent treated control

f Fisher's exact test versus diluent treated control

f Fisher's exact test versus diluent treated control

s Student's t-test versus diluent treated control

p = 0.037

s Student's t-test versus diluent treated control

p = 0.011

Results of Testing AVS-5587 Against Japanese Encephalitis Virus Challenge

Table 49

JE Experiment No. 41

	No. Dead/	No. Dead/			
Treatment	No. Uninfected	No. Infected	<u>p</u> a	$ADD + 1 SD^b$	pc
Untreated	0/5	10/10		13.4 <u>+</u> 2.1	
0.4% CMC	0/5	10/10	••	13.5 <u>+</u> 1.1	
AVS-5587					
200 mg/kg	0/5	1/10	0.00006	17.0 <u>+</u> 0.0	0.013
100 mg/kg	0/5	2/10	0.00036	20.0 ± 1.4	0.00002
50 mg/kg	0/5	1/10	0.00006	18.0 <u>+</u> 0.0	0.003
10 mg/kg	0/5	9/10		14.2 <u>+</u> 1.3	0.20
5 mg/kg	0/5	10/10		13.9 <u>+</u> 1.8	0.55
1 mg/kg	0/5	10/10		14.0 <u>+</u> 1.8	0.45
0.5 mg/kg	0/5	7/10	0.11	14.4 <u>+</u> 1.6	0.17
0.1 mg/kg	0.5	9/10	0.5	13.1 ± 2.1	0.61

p value resulting from Fisher's Exact test comparison of the mortality rates of drug-treated, virus-infected to diluent-treated, virus-infected control mice.

b ADD ± 1 SD = Average Day of Death ± 1 Standard Deviation.

p value resulting from Student's t-test comparison of the ADD of drug-treated, virus-infected to diluent-treated, virus-infected control mice.

4.3.4. Vaccinia Virus (VV):

A total of 9 compounds were assayed for activity against VV in vivo. These compounds are listed in Table 50. Four of the compounds were assessed for activity in the intracranial challenge model. These included AVS 1752 (ara-A; positive control drug), 1985, 1986 and 1987. Of these, only AVS 1752 had demonstrable activity against the virus induced mortality. All 9 compounds had some degree of activity against virus-induced tailpox lesions.

The cumulative results from these assays are presented in Table 51. The compound with the greatest activity other than the positive control drug (AVS 1752) was AVS 3679. In the first assay of this compound the activity was not marked. However, a second assay indicated significantly greater activity. This difference is believed to have resulted from changes in the drug preparation. In the second assay the compound was solubilized immediately prior to administration whereas in the first assay less consideration were given to drug stability.

Following AVS 3679 the greatest activity was seen with AVS 4225 then AVS 2875. Beyond these there were no marked differences in activity.

Since all of the compounds had some antiviral efficacy they should all be considered for additional dose, route and schedule studies. Additional information provided by pharmacokinetics studies would be beneficial in establishing the appropriate dose, route and schedule for maximal therapeutic efficacy. Consideration should be given to trying constant drug administration via Alzet[®] pumps or repeated daily dosing. The oral bioavailability should also be determined and the anti-vaccinia virus activity assessed using repeated daily oral dosing.

Table 50

Compounds Received for Testing Against In Vivo Vaccinia Virus

AVS#	IC Challenge Activity	Tailpox Activity	Maximum tolerated dose	Recommendation
1752 (Ara-A	A) yes	yes	>400 mg/kg/day	further studies
1985	no	yes	>293 mg/kg/day	further studies
1986	no	yes	250-500 mg/kg/day	further studies
1987	no	yes	>320 mg/kg/day	further studies
1988	ND	yes	300 mg/kg/day	further studies
2875	ND	yes	> 320 mg/kg/day	further studies
2994	ND	yes	300 mg/kg/day	further studies
3679	ND	yes	>300 mg/kg/day	further studies
4225	ND	yes	>300 mg/kg/day	further studies

ND - not done

Table 51

Activity of Selected AVS Compounds vs Vaccinia Virus (Tailpox Counts)

	% Reduction in	% Reduction in	
AVS#	Mean Tailpox Counts	Median Tailpox Cour	
1752			
300 mg/kg	58%	76%	
300 mg/kg	52%	83 %	
400 mg/kg	20%	47%	
400 mg/kg ¹	60%	88%	
300 mg/kg	77 %	88%	
300 mg/kg	46%	45%	
300 mg/kg	56%	38%	
400 mg/kg	73%	ND	
985			
293 mg/kg	55%	55%	
325 mg/kg	5%	44%	
325 mg/kg	37%	47%	
986			
250 mg/kg	39%	61%	
300 mg/kg	54%	86%	
100 mg/kg ¹	69 %	77 %	
987			
320 mg/kg	39%	51%	
988			
300 mg/kg	31%	64%	
875			
300 mg/kg	50%	50%	
994			
300 mg/kg	38%	52%	
225			
300 mg/kg	57%	67%	
679	100		
300 mg/kg	27%	45%	
300 mg/kg	97%	100%	
100 mg/kg ¹	55 %	67%	

¹Compounds administered subcutaneously rather than intraperitoneally.

5. **DISCUSSION**

Around 305 AVS compounds demonstrated antiviral activity at greater than 50% reduction levels against the Vaccinia Virus. Out of these, 34 AVS compounds appeared to have excellent *in vitro* antiviral potential with antiviral values that reached better than 95% reduction levels with TAI values that ranged from 11% to 83%. The best of these active leads should be studied further.

One hundred (100) compounds demonstrated *in vitro* antiviral activity at greater than or equal to 50% reduction levels against the Adenovirus. Some of the compounds found to be most effective against AD2 were: AVS-2296, 2700, 2980, 2986, 3593, 4070 and 4167.

We have performed 7654 in vitro antiviral assays against the YF Virus including the quality control tests. Around 306 mpounds demonstrated antiviral activity at greater than 50% reduction levels. Out of these assays, 02 AVS compounds appeared to have excellent in vitro antiviral potential with antiviral values that reached better than 95% reduction levels with TAI values that ranged from 5% to 99%. These results warrant that the best of these active leads would be studied further.

In this contract period, we performed 7873 in vitro antiviral assays against the JE Virus including quality control tests. Around 206 compounds demonstrated antiviral activity at greater than 50% reduction levels. Out of these assays, 45 AVS compounds appeared to have excellent in vitro antiviral potential with antiviral values that reached better than 95% reduction levels with TAI values that ranged from 9% to 95%. The results warrant that the best of these active leads would be studied further.

Against VE virus, 7319 in vitro antiviral assays were performed during this contract period including the quality control tests. Around 193 compounds demonstrated antiviral activity at greater than 50% reduction levels. Out of these assays, 15 AVS compounds appeared to have excellent in vitro antiviral potential with antiviral values that reached better than 95% reduction levels with TAI values that ranged from 7% to 87%. These results warrant that the best of these active leads would be studied further.

During this contract period, we performed 8221 in vitro antiviral assays against the PT virus including the quality control tests. Around 635 compounds demonstrated antiviral activity at greater than 50% reduction levels. Out of these assays, 107 AVS compounds appeared to have excellent in vitro antiviral potential with antiviral values that reached better than 95% reduction levels with TAI values that ranged from 10% to 96%. These results definitely warrant that the best out of these active leads would be studied further.

We performed 7833 in vitro assays against the SF virus including the quality control tests. Around 566 compounds demonstrated antiviral activity at greater than 50% reduction levels. Out of these assays, 95 AVS compounds appeared to have excellent in vitro antiviral potential with antiviral values that reached better than 95% reduction levels with TAI values that ranged from 5 - 97%. These results warrant that the best of these active leads would be studied further.

We have performed 2276 in vitro antiviral assays against the Pichinde Virus including the positive control tests. Around 197 compounds demonstrated antiviral activity at greater than 50% reduction levels and therapeutic indices of > 1.0. Five compounds (AVS-0646, 0140, 0148, 2350 and 3189) produced therapeutic indices of > 100. The results warrant that the best of these active leads would be studied further.

The prescreen protocol has successfully identified potential active materials (~5%) for further confirmatory testing. Confirmatory testing of these potential active compounds were carried out in the primary screen against a broader range of more virulent viruses (VV, YF, JE, VE, PT or SF). Sixty-seven percent (515/767) of the prescreen compounds showed some degree of activity against one or more of these more virulent viruses.

During the time that we conducted primary HIV testing on this contract, 14 AVS compounds produced anti-HIV activity with Therapeutic Indices that ranged from > 54 to > 1000. One NCI compound, NSC 614846 demonstrated significant activity comparable to the positive control drug, ddC. Several compounds showed confirmed anti-HIV activity versus the Feline and Simian Viruses (AVS-0001, 999, 2285 and 2639). AVS-0001 and AVS-2639 were active against the Murine Virus (MAIDS).

From the active *in vitro* lead compounds, the following have been advanced to appropriate *in vivo* animal model studies with the following results:

A total of 19 compounds were assessed for efficacy against Pichinde Virus in hamsters. Out of the compounds evaluated, only 2 (AVS-0001 and AVS-0206) had significant activity against Pichinde Virus in vivo. Both compounds demonstrated good potential for treatment of arenaviral infections.

None of the six compounds that we received to be tested against VE in mice had significant antiviral activity. However, five of the six compounds were not available in sufficient amounts for testing at maximum tolerated dose levels. Thus, it is possible that these compounds may show activity against VE if higher dose levels are tested.

A total of 13 compounds were received for testing against JE in mice. Out of these compounds, 7 had demonstrable activity against JE in vivo. These compounds were AVS-360, 361, 2811, 2812, 2979, 2980 and 5587. Of all the compounds demonstrating anti-JE activity, AVS-5587 may be the most desirable as the window between toxicity and efficacy is broader than that seen with the other compounds. Further studies of this compound and its analogs are strongly recommended.

Nine compounds were assayed for activity against VV in mice. Four of the compounds were assessed for activity in the intracranial challenge model. These included AVS-1752 (Ara-A; positive control drug), 1985, 1986 and 1987. Of these, only AVS-1752 had demonstrable activity against the virus induced mortality. All 9 compounds had some degree of activity against virus-induced tailpox lesions. The compound with the greatest activity other than the positive control drug (AVS-1752) was AVS-3679. Following AVS-3679 the greatest activity was seen with AVS-4225 then AVS-2875. Beyond these there were no marked differences in activity.

6. ACKNOWLEDGMENTS

Dr. Louis E. Holland II served as Assistant Program Manager for the exotic RNA virus screen. Dr. Gary J. Williams served as Task Leader for the SF and PIC viruses. Dr. Lorraine V. Brando served as Task Leader for the YF, JE, VEE, PT, FeLV, SAIDS viruses.

Mr. David Baggett (Associate Biologist), Ms. Jean Bailey (Assistant Biologist), Ms. Connie Bryant (Assistant Biologist), Ms. Sheri Campbell (Biological Technician), Ms. Sharon Chynoweth (Assistant Biologist), Ms. Joan Conway (Associate Biologist), Mr. Ali Danner (Biological Technician), Mr. Fred Davidson (Assistant Biologist), Ms. Lindsey Deckard (Assistant Microbiologist), Ms. Cynthia Doggett (Associate Biologist), Ms. Carol Eldridge (Associate Biologist), Mr. James Gallaspy (Assistant Biologist), Ms. Allison Heald (Assistant Biologist), Mr. Darryl Hicks (Assistant Biologist), Ms. Diane Horton (Associate Biologist), Mr. John Hultquist (Assistant Biologist), Ms. Jody Jones (Assistant Biologist), Ms. Karen Shelton Keith (Assistant Biologist), Mr. Richard Kirkman (Associate Microbiologist), Ms. Sandra Kooyer (Assistant Biologist), Ms. Edith Mayomi (Assistant Biologist), Ms. Sarah Pickett (Biological Research Technician), Mr. Daniel C. Potts (Assistant Biologist), Ms. Beverly Roberts (Associate Biologist), Ms. Rose Smith (Assistant Biologist), Ms. Ellen Stringfellow (Assistant Biologist), Mr. Robert Tubbs (Biological Technician), performed the *in vitro* antiviral evaluations against the DNA viruses and exotic RNA viruses.

Ms. Elizabeth A. Dulmadge (Research Biologist) supervised the day to day operations of the Centralized Cell Culture and Drug Preparation Laboratories.

Ms. Carrie Edwards (Biological Technician), Ms. LaJuana Farris (Biological Technician), Ms. Geraldine Jefferson (Assistant Biologist), Mr. Byron Lambert (Assistant Biologist), Ms. Barbara Toyer (Research Biologist) and Mr. Frank Vance (Biological Technician) performed the *in vivo* antiviral evaluations.

Dr. George C. Lavelle served as Assistant Program Manager for the anti-HIV screen and confirmatory testing. Dr. Jasbir B. Kahlon served as Task Leader for the HIV laboratory.

Mr. Donald Decker (Assistant Biologist), Mr. Joseph Johnson (Assistant Biologist), Mr. Jeffrey McCurdy (Associate Microbiologist), Ms. Teresa McDuffie (Assistant Biologist), Mr. Christopher McGee (Associate Microbiologist), Ms. Pamela Pruett (Assistant Biologist), Ms. Jeanine Qualls (Assistant Biologist), Mr. Thomas Rowe (Assistant Molecular Biologist), and Ms. Robin Worsham (Assistant Biologist) performed the primary *in vitro* antiviral evaluations against HIV. Ms. Bonnie Bowdon (Staff Biochemist), performed the HIV-immunofluorescence assays. Ms. Lucile White (Staff Biochemist) and Mr. James Konzelman (Assistant Chemist), performed the reverse transcriptase assays.

Ms. Joan Johnson (Assistant Statistician) and Ms. Santosh Niwas (Assistant Statistician) performed statistical analysis on the data. Ms. Kimberly Page (Biological Technician), Ms. Judith Talley (Biological Technician), Ms. Rose Vizzinia (Data Technician) helped with data processing and reporting.

Ms. Renee McCurdy (Assistant Programmer) designed and wrote several R-Base programs for in vitro data input and reporting. Mr. Martin Schulman and Mr. Steve Wideman (Programming Consultants) wrote the computer programs that enabled the MTT data to be automated from the plate reader in the laboratory to the printing of the "Antiviral MTT Assay" report (primary and prescreen) for submission to the sponsor.

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8. ABSTRACTS/PUBLICATIONS

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